

09/690,353

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NEWS 1 Web Page URLs for STN Seminar Schedule - N. America  
NEWS 2 Dec 17 The CA Lexicon available in the CAPLUS and CA files  
NEWS 3 Feb 06 Engineering Information Encompass files have new names  
NEWS 4 Feb 16 TOXLINE no longer being updated  
NEWS 5 Apr 23 Search Derwent WPINDEX by chemical structure  
NEWS 6 Apr 23 PRE-1967 REFERENCES NOW SEARCHABLE IN CAPLUS AND CA  
NEWS 7 May 07 DGENE Reload  
NEWS 8 Jun 20 Published patent applications (A1) are now in USPATFULL  
NEWS 9 JUL 13 New SDI alert frequency now available in Derwent's  
DWPI and DPCI  
NEWS 10 Aug 23 In-process records and more frequent updates now in  
MEDLINE  
NEWS 11 Aug 23 PAGE IMAGES FOR 1947-1966 RECORDS IN CAPLUS AND CA  
NEWS 12 Aug 23 Adis Newsletters (ADISNEWS) now available on STN  
NEWS 13 Sep 17 IMSworld Pharmaceutical Company Directory name change  
to PHARMASEARCH  
NEWS 14 Oct 09 Korean abstracts now included in Derwent World Patents  
Index  
NEWS 15 Oct 09 Number of Derwent World Patents Index updates increased  
NEWS 16 Oct 15 Calculated properties now in the REGISTRY/ZREGISTRY File  
NEWS 17 Oct 22 Over 1 million reactions added to CASREACT  
NEWS 18 Oct 22 DGENE GETSIM has been improved  
NEWS 19 Oct 29 AAASD no longer available  
  
NEWS EXPRESS August 15 CURRENT WINDOWS VERSION IS V6.0c,  
CURRENT MACINTOSH VERSION IS V6.0 (ENG) AND V6.0J (JP),  
AND CURRENT DISCOVER FILE IS DATED 07 AUGUST 2001  
NEWS HOURS STN Operating Hours Plus Help Desk Availability  
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NEWS PHONE Direct Dial and Telecommunication Network Access to STN  
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COST IN U.S. DOLLARS

FULL ESTIMATED COST

SINCE FILE	TOTAL
ENTRY	SESSION
0.30	0.30

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FILE 'REGISTRY' ENTERED AT 14:30:55 ON 02 NOV 2001  
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STRUCTURE FILE UPDATES: 1 NOV 2001 HIGHEST RN 366444-38-4  
DICTIONARY FILE UPDATES: 1 NOV 2001 HIGHEST RN 366444-38-4

TSCA INFORMATION NOW CURRENT THROUGH July 7, 2001

Please note that search-term pricing does apply when  
conducting SmartSELECT searches.

Crossover limits have been increased. See HELP CROSSOVER see  
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Calculated physical property data is now available. See HELP PROPERTIES  
for more information. See STNote 27, Searching Properties in the CAS  
Registry File, for complete details:  
<http://www.cas.org/ONLINE/STN/STNOTES/stnotes27.pdf>

=> e cobalamin?

E1	2	COBALACTONE/BI
E2	343	COBALAMIN/BI
E3	0 -->	COBALAMIN?/BI
E4	21	COBALAMINE/BI
E5	1	COBALAMINELACTAM/BI
E6	1	COBALAMINMONO/BI
E7	1	COBALAMINMONOCARBOXYL/BI
E8	1	COBALAMINMONOCARBOXYLIC/BI
E9	2	COBALAMINSULFONIC/BI
E10	1	COBALAMINTRI/BI
E11	1	COBALAMINTRICARBOXYL/BI
E12	1	COBALAMINTRICARBOXYLIC/BI

=> s e2

L1 343 COBALAMIN/BI

=> d

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LI ANSWER 1 OF 343 REGISTRY COPYRIGHT 2001 ACS  
RN 357492-04-7 REGISTRY  
CN 269Aa long cobalamin biosynthesis protein cobD (Sulfolobus tokodaii  
strain 7 gene ST2342) (9CI) (CA INDEX NAME)  
OTHER NAMES:  
CN GenBank AF000989-derived protein GI 15623463  
FS PROTEIN SEQUENCE  
MP Unspecified  
CI MAN  
SR CA  
LC STN Files: CA, CAPLUS

\*\*\* STRUCTURE DIAGRAM IS NOT AVAILABLE \*\*\*  
\*\*\* USE 'SQD' OR 'SQIDE' FORMATS TO DISPLAY SEQUENCE \*\*\*  
1 REFERENCES IN FILE CA (1967 TO DATE)  
1 REFERENCES IN FILE CAPLUS (1967 TO DATE)

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=> fil .search

COST IN U.S. DOLLARS

SINCE FILE

TOTAL

ENTRY

SESSION

FULL ESTIMATED COST

5.61

5.91

FILE 'MEDLINE' ENTERED AT 14:31:28 ON 02 NOV 2001

FILE 'CAPLUS' ENTERED AT 14:31:28 ON 02 NOV 2001

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FILE 'USPATFULL' ENTERED AT 14:31:28 ON 02 NOV 2001

CA INDEXING COPYRIGHT (C) 2001 AMERICAN CHEMICAL SOCIETY (ACS)

FILE 'EMBASE' ENTERED AT 14:31:28 ON 02 NOV 2001

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=> s l1

L2 50399 L1

<-----User Break----->

u

SEARCH ENDED BY USER

=> s l2 and (peptide? or polypeptide? or amino(w)acid?)

2 FILES SEARCHED...

4 FILES SEARCHED...

L3 3794 L2 AND (PEPTIDE? OR POLYPEPTIDE? OR AMINO(W) ACID?)

=> s l3 and (radionuclide? or radiolabel? or radioactiv? or radioisotop?)

L4 208 L3 AND (RADIONUCLIDE? OR RADIOLABEL? OR RADIOACTIV? OR RADIOISO  
TOP?)

=> s l4 and (chelate? or ligand?)

L5 72 L4 AND (CHELAT? OR LIGAND?)

=> dup rem l5

PROCESSING COMPLETED FOR L5

L6 68 DUP REM L5 (4 DUPLICATES REMOVED)

=> d ibib ab 1-

YOU HAVE REQUESTED DATA FROM 68 ANSWERS - CONTINUE? Y/(N):y

09/690,353

L6 ANSWER 1 OF 68 CAPLUS COPYRIGHT 2001 ACS  
 ACCESSION NUMBER: 2001:300552 CAPLUS  
 DOCUMENT NUMBER: 134:323118  
 TITLE: Cobalamin conjugates useful as tumor imaging and therapeutic agents  
 INVENTOR(S): Hogenkamp, Henricus P. C.; Collins, Douglas A.  
 PATENT ASSIGNEE(S): Mayo Foundation for Medical Education and Research, USA; Regents of the University of Minnesota  
 SOURCE: PCT Int. Appl., 46 pp.  
 DOCUMENT TYPE: Patent  
 LANGUAGE: English  
 FAMILY ACC. NUM. COUNT: 1  
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2001028595	A1	20010426	WO 2000-US10098	20000415
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, ME, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GN, GW, ML, MR, NE, SN, TD, TG				

PRIORITY APPL. INFO.: US 1999-159753 A2 19991015  
 OTHER SOURCE(S): MARPAT 134:323118  
 AB The invention provides detectably labeled cobalamin deriva. which are useful for medical treatment and diagnosis. Cobalamin conjugates have peptides or amino acids linked to metal radionuclide chelates or nonmetallic radionuclides.  
 REFERENCE COUNT: 4  
 REFERENCE(S): (1) Biotech Australia Pty Ltd; WO 9427613 A 1994 CAPLUS  
 (2) Collins, D; US 5739313 A 1998 CAPLUS  
 (3) Collins, D; WO 0062808 A 2000 CAPLUS  
 (4) Grissom, C; WO 9808859 A 1998 CAPLUS

L6 ANSWER 2 OF 68 USPATFULL  
 ACCESSION NUMBER: 2001:163361 USPATFULL  
 TITLE: Discrete-length polyethylene glycols  
 INVENTOR(S): Wilbur, D. Scott, Edmonds, WA, United States  
 Patthare, Pradip M., Seattle, WA, United States  
 PATENT ASSIGNEE(S): The University of Washington, Seattle, WA, United States (U.S. corporation)

NUMBER	KIND	DATE
US 6294697	B1	20010925
US 1998-62286		19980417 (9)

Continuation of Ser. No. WO 1996-US16760, filed on 18 Oct 1996

NUMBER	DATE
US 1995-5599	19951019 (60)

PRIORITY INFORMATION: US 1995-5599 19951019 (60)  
 DOCUMENT TYPE: Utility  
 FILE SEGMENT: GRANTED  
 PRIMARY EXAMINER: Padmanabhan, Sreeni  
 LEGAL REPRESENTATIVE: Christensen O'Connor Johnson & Kindness PLLC  
 NUMBER OF CLAIMS: 38  
 EXEMPLARY CLAIM: 1  
 NUMBER OF DRAWINGS: 14 Drawing Figure(s); 14 Drawing Page(s)  
 LINE COUNT: 1072  
 CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The present invention provides discrete-length polyethylene glycol and polyethylene glycol containing compounds and methods for their preparation.

L6 ANSWER 3 OF 68 USPATFULL  
 ACCESSION NUMBER: 2001:152712 USPATFULL  
 TITLE: nrdg  
 INVENTOR(S): Black, Michael Terence, Chester Springs, PA, United States  
 PATENT ASSIGNEE(S): SmithKline Beecham Corporation, Philadelphia, PA, United States (U.S. corporation)

NUMBER	KIND	DATE
US 6287804	B1	20010911
US 1998-40213		19980317 (9)

PATENT INFORMATION: US 6287804 B1 20010911  
 APPLICATION INFO.: US 1998-40213 19980317 (9)  
 DOCUMENT TYPE: Utility  
 FILE SEGMENT: GRANTED  
 PRIMARY EXAMINER: Martinell, James  
 LEGAL REPRESENTATIVE: Gimmi, Edward R., Deibert, Thomas S., King, William T.  
 NUMBER OF CLAIMS: 13  
 EXEMPLARY CLAIM: 1  
 LINE COUNT: 2108  
 CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The invention provides nrdg polypeptides and polynucleotides encoding nrdg polypeptides and methods for producing such polypeptides by recombinant techniques. Also provided are methods for utilizing nrdg polypeptides to screen for antibacterial compounds.

L6 ANSWER 4 OF 68 USPATFULL  
 ACCESSION NUMBER: 2001:131279 USPATFULL  
 TITLE: Compositions of cobalamin and related corrinoids, and uses thereof  
 INVENTOR(S): Sarill, William J., 78 Hibbert St., Arlington, MA, United States 02154  
 Brennan, Thomas F., 44 A Gail Dr., Nyack, NY, United States 10960

NUMBER	KIND	DATE
US 6274564	B1	20010814
US 1997-936781		19970917 (8)

PATENT INFORMATION: US 6274564 B1 20010814  
 APPLICATION INFO.: US 1997-936781 19970917 (8)

NUMBER	DATE
US 1996-25298	19960918 (60)
US 1997-41750	19970328 (60)

PRIORITY INFORMATION: US 1996-25298 19960918 (60)  
 US 1997-41750 19970328 (60)  
 DOCUMENT TYPE: Utility  
 FILE SEGMENT: GRANTED  
 PRIMARY EXAMINER: Travers, Russell  
 LEGAL REPRESENTATIVE: Lahive & Cockfield, LLP  
 NUMBER OF CLAIMS: 17  
 EXEMPLARY CLAIM: 1  
 LINE COUNT: 1907  
 CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB Novel compositions cobalamin and related corrinoids, and uses thereof, are disclosed. The novel compositions include a corrin, a first amino acid having a side chain which includes a basic or positively charged moiety; and a second amino acid with an uncharged side chain which includes at least one heteroatom.  
 The compositions are useful for, inter alia, treatment of cobalamin deficiency.

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L6 ANSWER 5 OF 68 USPATFULL  
 ACCESSION NUMBER: 2001:98064 USPATFULL  
 TITLE: Ribb  
 INVENTOR(S): Black, Michael Terance, Chester Springs, PA, United States  
 Shilling, Lisa Kathleen, Newtown, PA, United States  
 Stodola, Robert King, Flourtown, PA, United States  
 Warren, Richard Lloyd, Blue Bell, PA, United States  
 Kosmatka, Anna Lisa, Doylestown, PA, United States  
 Nicholas, Richard Oakley, Collegeville, PA, United States  
 Palmer, Leslie Marie, Audubon, PA, United States  
 Lonetto, Michael Arthur, Collegeville, PA, United States  
 Fedon, Jason Craig, Strafford, PA, United States  
 Hodgson, John Edward, Malvern, PA, United States  
 Knowles, David Justin Charles, Boroughbridge, United Kingdom  
 PATENT ASSIGNEE(S): SmithKline Beecham Corporation, Philadelphia, PA, United States (U.S. corporation)  
 SmithKline Beecham plc, United Kingdom (non-U.S. corporation)

NUMBER	KIND	DATE
US 6252044	B1	20010626
US 1997-977555		19971125 (8)

NUMBER	DATE
US 1996-24022	19960816 (60)

PATENT INFORMATION: US 1996-24022  
 DOCUMENT TYPE: Utility  
 FILE SEGMENT: GRANTED  
 PRIMARY EXAMINER: Minnifield, Nita  
 LEGAL REPRESENTATIVE: Gimmi, Edward R., Diebert, Thomas S., King, William T.  
 NUMBER OF CLAIMS: 16  
 EXEMPLARY CLAIM: 1  
 LINE COUNT: 1599

CAS INDEXING IS AVAILABLE FOR THIS PATENT.  
 AB The invention provides ribb polypeptides and polynucleotides encoding ribb polypeptides and methods for producing such polypeptides by recombinant techniques. Also provided are methods for utilizing ribb polypeptides to screen for antibacterial compounds.

L6 ANSWER 7 OF 68 USPATFULL  
 ACCESSION NUMBER: 2001:17971 USPATFULL  
 TITLE: Transcobalamin mediated transport of vitamins B12 in intrinsic factor or receptor deficient patient  
 INVENTOR(S): Seetharam, Bellur, Brookfield, WI, United States  
 Bose, Santanu, San Francisco, CA, United States  
 NOW Research Foundation, Milwaukee, WI, United States (U.S. corporation)

NUMBER	KIND	DATE
US 6183723	B1	20010206
US 1998-9995		19980121 (9)

PATENT INFORMATION: US 6183723  
 APPLICATION INFO.: US 1998-9995  
 DOCUMENT TYPE: Utility  
 FILE SEGMENT: Granted  
 PRIMARY EXAMINER: Saucier, Sandra E.  
 ASSISTANT EXAMINER: Afrenova, Vera  
 LEGAL REPRESENTATIVE: Quarles & Brady LLP  
 NUMBER OF CLAIMS: 1  
 EXEMPLARY CLAIM: 1  
 NUMBER OF DRAWINGS: 7 Drawing Figure(s); 6 Drawing Page(s)  
 LINE COUNT: 869

CAS INDEXING IS AVAILABLE FOR THIS PATENT.  
 AB A method for oral treatment of patients using different types of drugs that are not usually transported to circulation, if administered orally, is disclosed. Because the TC II-Cbl complex is stable to acid and proteolytic enzymes both outside and inside the intestinal absorptive cells, the method consists of oral administration of a drug bound to TC II-Cbl complex. In addition, the method can also be used for delivering Cbl to a large number of patients who do not absorb Cbl due to a variety of causes such as surgery of their stomach (ulcers) or of their terminal ileum (Crohn's disease).

L6 ANSWER 6 OF 68 USPATFULL  
 ACCESSION NUMBER: 2001:97419 USPATFULL  
 TITLE: Method and a system for enhanced in vivo clearance of diagnostic and/or therapeutic agents by extracorporeal depletion, and the use of said agents for said purpose  
 INVENTOR(S): Nilsson, Rune, Lund, Sweden  
 Lindgren, Lars, Lund, Sweden  
 Norrgren, Kristina, ANG.karp, Sweden  
 Sandberg, Bengt, Lund, Sweden  
 Sjogren, Hans Olof, Lund, Sweden  
 Strand, Sven-Erik, Lund, Sweden  
 PATENT ASSIGNEE(S): Mitra Medical Technology AB, Lund, Sweden (non-U.S. corporation)

NUMBER	KIND	DATE
US 6251394	B1	20010626
WO 9212730		19920806
US 1993-90047		19931012 (8)
WO 1992-SE20		19920115
		19931012 PCT 371 date
		19931012 PCT 102(e) date

NUMBER	DATE
SE 1991-142	19910117

PATENT INFORMATION: SE 1991-142  
 DOCUMENT TYPE: Utility  
 FILE SEGMENT: GRANTED  
 PRIMARY EXAMINER: Witz, Jean C.  
 LEGAL REPRESENTATIVE: Swanson & Bratschun L  
 NUMBER OF CLAIMS: 3  
 EXEMPLARY CLAIM: 1  
 NUMBER OF DRAWINGS: 4 Drawing Figure(s); 2 Drawing Page(s)  
 LINE COUNT: 656

CAS INDEXING IS AVAILABLE FOR THIS PATENT.  
 AB A method and a system is described for reducing non-target levels of specific molecules intended for diagnostic and/or therapeutic applications to vertebrate hosts, wherein said molecules are administered to a vertebrate host and kept therein for a certain time in order to be concentrate to the target by being attached thereto. The molecules which are not attached to the target are removed from the blood circulation system or at least reduced to a lower concentration by passing the blood through an extra-corporeal device.

L6 ANSWER 8 OF 68 USPATFULL  
 ACCESSION NUMBER: 2001:4509 USPATFULL  
 TITLE: ribb  
 INVENTOR(S): Black, Michael Terance, Chester Springs, PA, United States  
 Fedon, Jason Craig, Strafford, PA, United States  
 Hodgson, John Edward, Malvern, PA, United States  
 Knowles, David Justin Charles, Boroughbridge, United Kingdom  
 Lonetto, Michael Arthur, Collegeville, PA, United States  
 Kosmatka, Anna Lisa, Doylestown, PA, United States  
 Nicholas, Richard Oakley, Collegeville, PA, United States  
 Palmer, Leslie Marie, Audubon, PA, United States  
 Shilling, Lisa Kathleen, Newtown, PA, United States  
 Stodola, Robert King, Flourtown, PA, United States  
 Warren, Richard Lloyd, Blue Bell, PA, United States  
 SmithKline Beecham Corporation, Philadelphia, PA, United States (U.S. corporation)  
 SmithKline Beecham plc, United Kingdom (non-U.S. corporation)

NUMBER	KIND	DATE
US 6171835	B1	20010109
US 1999-385288		19990830 (9)

PATENT INFORMATION: US 6171835  
 APPLICATION INFO.: US 1999-385288  
 RELATED APPL. INFO.: Division of Ser. No. US 1997-378454, filed on 25 Nov 1997, now patented, Pat. No. US 6017728 Continuation of Ser. No. US 1997-911503, filed on 15 Aug 1997  
 Continuation of Ser. No. WO 1997-US14436, filed on 15 Aug 1997

NUMBER	DATE
US 1996-24022	19960816 (60)

PATENT INFORMATION: US 1996-24022  
 DOCUMENT TYPE: Patent  
 FILE SEGMENT: Granted  
 PRIMARY EXAMINER: Carlson, Karen Cochrane  
 LEGAL REPRESENTATIVE: Gimmi, Edward R., Diebert, Thomas S., King, William T.  
 NUMBER OF CLAIMS: 14  
 EXEMPLARY CLAIM: 1  
 LINE COUNT: 1355

CAS INDEXING IS AVAILABLE FOR THIS PATENT.  
 AB The invention provides ribb polypeptides and polynucleotides encoding ribb polypeptides and methods for producing such polypeptides by recombinant techniques. Also provided are methods for utilizing ribb polypeptides to screen for antibacterial compounds.

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L6 ANSWER 9 OF 68 USPATFULL  
 ACCESSION NUMBER: 2000:157392 USPATFULL  
 TITLE: Vitamin B.sub.12 derivatives and methods for their preparation  
 INVENTOR(S): Russell-Jones, Gregory J., New South Wales, Australia  
 McEwan, John F., New South Wales, Australia  
 PATENT ASSIGNEE(S): Biotech Australia Pty Limited, Roseville, Australia (non-U.S. corporation)

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 6150341		20001121
APPLICATION INFO.:	US 1999-330167		19990611 (9)

	NUMBER	DATE
PRIORITY INFORMATION:	AU 1998-4050	19980612
DOCUMENT TYPE:	Utility	
FILE SEGMENT:	Granted	
PRIMARY EXAMINER:	Wilson, James O.	
LEGAL REPRESENTATIVE:	Foley & Lardner	
NUMBER OF CLAIMS:	26	
EXEMPLARY CLAIM:	1	
NUMBER OF DRAWINGS:	1 Drawing Figure(s); 1 Drawing Page(s)	
LINE COUNT:	760	

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB This invention relates to methods for preparing vitamin B.sub.12 (VB.sub.12) derivatives suitable for linking to a polymer, nanoparticle or therapeutic agent, protein or peptide. The methods involve reacting the 5'-OH group of VB.sub.12 or an analogue thereof with an active carbonyl electrophile and subsequently obtaining said VB.sub.12 derivatives. The invention also relates to novel VB.sub.12 derivatives, VB.sub.12 derivatives prepared by the methods of the present invention and uses thereof in the preparation of in the preparation of polymer complexes or nanoparticles.

L6 ANSWER 10 OF 68 USPATFULL  
 ACCESSION NUMBER: 2000:125190 USPATFULL  
 TITLE: pcrA protein from Streptococcus pneumoniae  
 INVENTOR(S): Black, Michael Terence, Chester Springs, PA, United States  
 Hodgeon, John Edward, Malvern, PA, United States  
 Knowles, David Justin Charles, Redhill, United Kingdom  
 Lonetto, Michael Arthur, Collegeville, PA, United States  
 Nicholas, Richard O., Collegeville, PA, United States  
 Stodola, Robert King, Flourtown, PA, United States  
 Holmes, David J., West Chester, PA, United States  
 PATENT ASSIGNEE(S): SmithKline Beecham Corporation, Philadelphia, PA, United States (U.S. corporation)

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 6121414		20000919
APPLICATION INFO.:	US 1998-185825		19981104 (9)
RELATED APPLN. INFO.:	Division of Ser. No. US 1997-889711, filed on 8 Jul 1997, now patented, Pat. No. US 5858718		

	NUMBER	DATE
DOCUMENT TYPE:	Utility	
FILE SEGMENT:	Granted	
PRIMARY EXAMINER:	Duffy, Patricia A.	
LEGAL REPRESENTATIVE:	Gimmi, Edward R., Deibert, Thomas S., King, William T.	
NUMBER OF CLAIMS:	16	
EXEMPLARY CLAIM:	1	
LINE COUNT:	1638	

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The invention provides pcrA polypeptides and DNA (RNA) encoding pcrA polypeptides and methods for producing such polypeptides by recombinant techniques. Also provided are methods for utilizing pcrA polypeptides to screen for antibacterial compounds.

L6 ANSWER 11 OF 68 USPATFULL  
 ACCESSION NUMBER: 2000:84267 USPATFULL  
 TITLE: Water soluble vitamin B.sub.12 receptor modulating agents and methods related thereto  
 INVENTOR(S): Morgan, Jr., A. Charles, Mill Creek, WA, United States  
 Wilbur, D. Scott, Edmonds, WA, United States  
 Patbare, Pradip M., Seattle, WA, United States  
 PATENT ASSIGNEE(S): The University of Washington, Seattle, WA, United States (U.S. corporation)  
 Receptagen Corporation, Edmonds, WA, United States (U.S. corporation)

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 6083926		20000704
APPLICATION INFO.:	US 1998-200422		19981123 (9)
RELATED APPLN. INFO.:	Division of Ser. No. US 1995-545151, filed on 19 Oct 1995, now patented, Pat. No. US 5840712 which is a continuation-in-part of Ser. No. WO 1995-US4404, filed on 7 Apr 1995 which is a continuation-in-part of Ser. No. US 1995-406191, filed on 16 Mar 1995, now patented, Pat. No. US 5840880 which is a continuation-in-part of Ser. No. US 1995-406192, filed on 16 Mar 1995, now patented, Pat. No. US 5739287 and a continuation-in-part of Ser. No. US 1995-406194, filed on 16 Mar 1995, now patented, Pat. No. US 5869465		

which is a continuation-in-part of Ser. No. US 1994-224831, filed on 8 Apr 1994, now abandoned

	NUMBER	DATE
DOCUMENT TYPE:	Utility	
FILE SEGMENT:	Granted	
PRIMARY EXAMINER:	Fonda, Kathleen K.	
LEGAL REPRESENTATIVE:	Seed Intellectual Property Law Group PLLC	
NUMBER OF CLAIMS:	16	
EXEMPLARY CLAIM:	1	
NUMBER OF DRAWINGS:	28 Drawing Figure(s); 18 Drawing Page(s)	
LINE COUNT:	3274	

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB Vitamin B.sub.12 receptor modulating agents capable of modulating cell surface receptors by affecting the cell surface receptor trafficking pathway are disclosed. The vitamin B.sub.12 receptor modulating agents are comprised of a covalently bound rerouting moiety and targeting moiety linked by a water-solubilizing linker.

L6 ANSWER 12 OF 68 USPATFULL  
 ACCESSION NUMBER: 2000:80599 USPATFULL  
 TITLE: Nucleophilic polysubstituted aryl acridinium ester conjugates and syntheses thereof  
 INVENTOR(S): Law, Say-Jong, Westwood, MA, United States  
 PATENT ASSIGNEE(S): Bayer Corporation, East Walpole, MA, United States (U.S. corporation)

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 6080591		20000627
APPLICATION INFO.:	US 1997-920372		19970829 (8)
RELATED APPLN. INFO.:	Continuation of Ser. No. US 1993-32947, filed on 17 Mar 1993, now patented, Pat. No. US 5663074 which is a continuation of Ser. No. US 1992-871601, filed on 17 Apr 1992, now patented, Pat. No. US 5241070 which is a continuation of Ser. No. US 1988-249620, filed on 26 Sep 1988, now abandoned		

	NUMBER	DATE
DOCUMENT TYPE:	Utility	
FILE SEGMENT:	Granted	
PRIMARY EXAMINER:	Wortman, Donna	
ASSISTANT EXAMINER:	Brumback, Brenda G.	
NUMBER OF CLAIMS:	15	
EXEMPLARY CLAIM:	1	
NUMBER OF DRAWINGS:	18 Drawing Figure(s); 15 Drawing Page(s)	
LINE COUNT:	1138	

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB This invention is directed to novel nucleophilic polysubstituted aryl acridinium conjugates and the methods for preparation thereof. The novel nucleophilic polysubstituted aryl acridinium conjugates are useful in biological assays, including novel assays for the determination of Vitamin B.sub.12, folate, cortisol, estradiol, and thromboxane B.sub.2.

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L6 ANSWER 13 OF 68 USPATFULL  
 ACCESSION NUMBER: 2000:18261 USPATFULL  
 TITLE: Glucose kinase from Streptococcus pneumoniae  
 INVENTOR(S): Black, Michael Terence, Chester Springs, PA, United States  
 Hodgson, John Edward, Malvern, PA, United States  
 Knowles, David Justin Charles, Redhill, United Kingdom  
 Lonetto, Michael Arthur, Collegeville, PA, United States  
 Nicholas, Richard O., Collegeville, PA, United States  
 Stodola, Robert King, Flourtown, PA, United States  
 Burnham, Martin Karl Russel, Norristown, PA, United States  
 PATENT ASSIGNEE(S): SmithKline Beecham Corporation, Philadelphia, PA, United States (U.S. corporation)  
 NUMBER KIND DATE  
 PATENT INFORMATION: US 6025175 20000215  
 APPLICATION INFO.: US 1998-110910 19980706 (9)  
 RELATED APPLN. INFO.: Division of Ser. No. US 1997-896083, filed on 17 Jul 1997, now patented, Pat. No. US 5840560  
 DOCUMENT TYPE: Utility  
 FILE SEGMENT: Granted  
 PRIMARY EXAMINER: Hobbs, Lisa J.  
 LEGAL REPRESENTATIVE: Gimmi, Edward R., King, William T., Deibert, Thomas S.  
 NUMBER OF CLAIMS: 18  
 EXEMPLARY CLAIM: 1  
 LINE COUNT: 1521  
 CAS INDEXING IS AVAILABLE FOR THIS PATENT.  
 AB The invention provides Glucose Kinase polypeptides and DNA (RNA) encoding Glucose Kinase polypeptides and methods for producing such polypeptides by recombinant techniques. Also provided are methods for utilizing Glucose Kinase polypeptides to screen for antibacterial compounds.

L6 ANSWER 14 OF 68 USPATFULL  
 ACCESSION NUMBER: 2000:9717 USPATFULL  
 TITLE: ribH  
 INVENTOR(S): Black, Michael Terence, Chester Springs, PA, United States  
 Fedon, Jason Craig, Strafford, PA, United States  
 Hodgson, John Edward, Malvern, PA, United States  
 Knowles, David Justin Charles, Boroughbridge, United Kingdom  
 Komanaka, Anna Lisa, Doylestown, PA, United States  
 Nicholas, Richard Oakley, Collegeville, PA, United States  
 Palmer, Leslie Marie, Audubon, PA, United States  
 Shilling, Lisa Kathleen, Newtown, PA, United States  
 Stodola, Robert King, Flourtown, PA, United States  
 Warren, Richard Lloyd, Blue Bell, PA, United States  
 Lonetto, Michael Arthur, Collegeville, PA, United States  
 PATENT ASSIGNEE(S): SmithKline Beecham Corporation, Philadelphia, PA, United States (U.S. corporation)  
 SmithKline Beecham plc, United Kingdom (non-U.S. corporation)  
 NUMBER KIND DATE  
 PATENT INFORMATION: US 6017728 20000125  
 APPLICATION INFO.: US 1997-978454 19971125 (8)  
 NUMBER DATE  
 PRIORITY INFORMATION: US 1996-24022 19960816 (60)  
 DOCUMENT TYPE: Utility  
 FILE SEGMENT: Granted  
 PRIMARY EXAMINER: Carlson, Karen Cochrane  
 LEGAL REPRESENTATIVE: Gimmi, Edward R., King, William T., Deibert, Thomas S.  
 NUMBER OF CLAIMS: 28  
 EXEMPLARY CLAIM: 1  
 LINE COUNT: 1624  
 CAS INDEXING IS AVAILABLE FOR THIS PATENT.  
 AB The invention provides ribH polypeptides and polynucleotides encoding ribH polypeptides and methods for producing such polypeptides by recombinant techniques. Also provided are methods for utilizing ribH polypeptides to screen for antibacterial compounds.

L6 ANSWER 15 OF 68 MEDLINE MEDLINE DUPLICATE 1  
 ACCESSION NUMBER: 2000442250  
 DOCUMENT NUMBER: 20443608 PubMed ID: 10990201  
 TITLE: Oral absorption of peptides through the cobalamin (vitamin B12) pathway in the rat intestine.  
 AUTHOR: Alsenz J; Russell-Jones G J; Westwood S; Levat-Trafit B; de Smidt P C  
 CORPORATE SOURCE: Preclinical Research Department, F. Hoffman-La Roche Ltd., Basle, Switzerland.. jochen.alsenz@roche.com  
 SOURCE: PHARMACEUTICAL RESEARCH, (2000 Jul) 17 (7) 825-32.  
 PUB. COUNTRY: United States  
 LANGUAGE: English  
 FILE SEGMENT: Priority Journals  
 ENTRY MONTH: 200101  
 ENTRY DATE: Entered STM: 20010322  
 Last Updated on STM: 20010322  
 Entered Medline: 20010118  
 AB PURPOSE: This study was aimed at examining the extent and mechanism of uptake of cobalamin (Cbl)-conjugated peptides in vitro and in vivo. METHODS: To enable acquisition of quantitative absorption data of Cbl-peptides, metabolically stable octapeptides (DP3), with (Cbl-Hex-DP3) or without a hexyl spacer (Cbl-DP3), were coupled to Cbl and radiolabeled. For comparison, LHRH coupled to Cbl was used as metabolically susceptible peptide. Biological recognition of Cbl-peptides was studied in the physiological order: binding by Intrinsic Factor (IF), recognition and transport of the IF-complexes by IF-Cbl receptors (IFCR) on Caco-2 monolayers and oral absorption of the Cbl-conjugates in the rat. RESULTS: All Cbl-peptides bound to IF and the IF-complexes were recognized by IFCR receptors on Caco-2 monolayers. Binding was saturable and could be inhibited by a 20-fold excess of IF-Cbl, but not of Non-intrinsic Factor (NIF)-Cbl. Oral administration of these ligands to rats resulted in absorption of 53%, 45%, 42%, and 23% of the applied radioactivity for Cbl, Cbl-LHRH, Cbl-Hex-DP3, and Cbl-DP3, respectively. Simultaneous administration of a >10(5)-fold excess of unlabeled Cbl reduced uptake of all compounds to <4%. Tissue distribution and elimination of the metabolically stable Cbl-conjugates were comparable to Cbl. CONCLUSIONS: The endogenous Cbl uptake pathway can be exploited for oral peptide delivery as indicated by the specific and high (40-45%) uptake of metabolically stable Cbl-coupled octapeptides.

L6 ANSWER 16 OF 68 USPATFULL  
 ACCESSION NUMBER: 1999:170440 USPATFULL  
 TITLE: Cell culturing method and medium  
 INVENTOR(S): Curcio, Francesco, Pagnacco, Italy  
 Cocchi, Hayden G., East Sebago, ME, United States  
 Ambesi-Impionato, F. Saverio, Udine, Italy  
 PATENT ASSIGNEE(S): Livercell L.L.C., East Sebago, ME, United States (U.S. corporation)  
 NUMBER KIND DATE  
 PATENT INFORMATION: US 6008047 19991228  
 APPLICATION INFO.: US 1998-66897 19980428 (9)  
 RELATED APPLN. INFO.: Continuation-in-part of Ser. No. US 1993-480022, filed on 7 Jun 1993, now patented, Pat. No. US 5888816 which is a continuation of Ser. No. US 1993-83772, filed on 30 Jun 1993, now abandoned which is a continuation-in-part of Ser. No. US 1993-44010, filed on 8 Apr 1993, now abandoned  
 DOCUMENT TYPE: Utility  
 FILE SEGMENT: Granted  
 PRIMARY EXAMINER: Lankford, Jr., Leon B.  
 ASSISTANT EXAMINER: Tate, Christopher R.  
 LEGAL REPRESENTATIVE: Bundock Jr., John P.  
 NUMBER OF CLAIMS: 16  
 EXEMPLARY CLAIM: 1  
 NUMBER OF DRAWINGS: 18 Drawing Figure(s); 11 Drawing Page(s)  
 LINE COUNT: 2290  
 CAS INDEXING IS AVAILABLE FOR THIS PATENT.  
 AB The present invention provides a method for producing an expanded non-transformed cell culture of human liver cells comprising the steps of: (1) preparing partially purified, minced human liver tissue, (2) concentrating the resulting cells and tissue pieces, (3) resuspending the concentrated tissue cells and pieces in a growth medium, (4) culturing the resuspended cells in the growth medium for a time and under conditions to effect sustained cell division, and (5) passaging the cultured human liver cells periodically to expand the culture. The growth medium comprises a combination of a basal medium and ingredients to provide a medium in which the cultured human liver cells are selectively proliferated without being transformed, providing an expanded culture of proliferated, functionally differentiated human liver cells that is substantially free of fibroblast, macrophage and capillary endothelial cells. Also provided is the improvement of harvesting cells of the expanded culture at a selected PDL preferably>5, providing a high density cell suspension of such proliferated human liver cells, and incubating such high density cell suspension in a calm-down medium to induce a mitotically quiescent state and, using a culture procedure which encourages aggregation, making the cells adhere tightly to form a three-dimensional cell organization typical of the organ of origin, thereby forming organoids.



09/690,353

L6 ANSWER 17 OF 68 USPATFULL  
 ACCESSION NUMBER: 1999:163707 USPATFULL  
 TITLE: Clioquinol for the treatment of Alzheimer's disease  
 INVENTOR(S): Gerolymatos, Panayotis N., Kryoneri Attikis, Greece  
 PATENT ASSIGNEE(S): P.N. Gerolymatos S.A., Kryoneri Attikis, Greece (non-U.S. corporation)

NUMBER	KIND	DATE
US 6001852		19991214
US 1998-23544		19980213 (9)

PATENT INFORMATION: US 6001852  
 APPLICATION INFO.: US 1998-23544  
 RELATED APPLN. INFO.: Continuation-in-part of Ser. No. WO 1997-1B983, filed on 8 Aug 1997

NUMBER	DATE
GR 1996-960100286	19960813

PRIORITY INFORMATION: GR 1996-960100286  
 DOCUMENT TYPE: Utility  
 FILE SEGMENT: Granted  
 PRIMARY EXAMINER: Criares, Theodore J.  
 LEGAL REPRESENTATIVE: Pennie & Edmonds LLP  
 NUMBER OF CLAIMS: 40  
 EXEMPLARY CLAIM: 1  
 NUMBER OF DRAWINGS: 4 Drawing Figure(s); 2 Drawing Page(s)  
 LINE COUNT: 1170  
 CAS INDEXING IS AVAILABLE FOR THIS PATENT.  
 AB A new pharmaceutical composition is disclosed that comprises clioquinol, vitamin B.sub.12, and, optionally, pharmaceutical acceptable carriers and/or excipients. The use of the pharmaceutical composition removes or alleviates the side effects of clioquinol.

L6 ANSWER 18 OF 68 USPATFULL  
 ACCESSION NUMBER: 1999:155706 USPATFULL  
 TITLE: Pharmaceutical compositions comprising clioquinol in combination with vitamin B12 and therapeutic and prophylactic uses thereof  
 INVENTOR(S): Gerolymatos, Panayotis N., Kryoneri Attikis, Greece  
 PATENT ASSIGNEE(S): P.N. Gerolymatos S.A., Kryoneri Attikis, Greece (non-U.S. corporation)

NUMBER	KIND	DATE
US 5994323		19991130
US 1998-23542		19980213 (9)

PATENT INFORMATION: US 5994323  
 APPLICATION INFO.: US 1998-23542

NUMBER	DATE
GR 1997-970100507	19971231

PRIORITY INFORMATION: GR 1997-970100507  
 DOCUMENT TYPE: Utility  
 FILE SEGMENT: Granted  
 PRIMARY EXAMINER: Criares, Theodore J.  
 LEGAL REPRESENTATIVE: Pennie & Edmonds LLP  
 NUMBER OF CLAIMS: 33  
 EXEMPLARY CLAIM: 1  
 NUMBER OF DRAWINGS: 4 Drawing Figure(s); 5 Drawing Page(s)  
 LINE COUNT: 1039  
 CAS INDEXING IS AVAILABLE FOR THIS PATENT.  
 AB A pharmaceutical composition is disclosed that comprises clioquinol, vitamin B.sub.12, and, optionally, pharmaceutical acceptable carriers and/or excipients. The use of the pharmaceutical composition removes or alleviates the side effects of clioquinol.

L6 ANSWER 19 OF 68 USPATFULL  
 ACCESSION NUMBER: 1999:155699 USPATFULL  
 TITLE: Method of preparing polynucleotide-carrier complexes for delivery to cells  
 INVENTOR(S): Lollo, Charles P., Encinitas, CA, United States  
 Mockler, Todd C., Carlsbad, CA, United States  
 Kwok, Deborah Y., Carlsbad, CA, United States  
 PATENT ASSIGNEE(S): The Immune Response Corporation, Carlsbad, CA, United States (U.S. corporation)

NUMBER	KIND	DATE
US 5994316		19991130
US 1998-604306		19960221 (8)

PATENT INFORMATION: US 5994316  
 APPLICATION INFO.: US 1998-604306  
 DOCUMENT TYPE: Utility  
 FILE SEGMENT: Granted  
 PRIMARY EXAMINER: Campbell, Bruce R.  
 LEGAL REPRESENTATIVE: Lahive & Cockfield, LLP, Remillard, Jane E.  
 NUMBER OF CLAIMS: 28  
 EXEMPLARY CLAIM: 1  
 NUMBER OF DRAWINGS: 16 Drawing Figure(s); 15 Drawing Page(s)  
 LINE COUNT: 1587  
 CAS INDEXING IS AVAILABLE FOR THIS PATENT.  
 AB An improved method of forming substantially disperse and homogeneous polynucleotide-carrier complexes is disclosed. The polynucleotide-carrier complexes can be administered in vivo to obtain significant levels and duration of gene expression.

L6 ANSWER 20 OF 68 USPATFULL  
 ACCESSION NUMBER: 1999:89277 USPATFULL  
 TITLE: riba  
 INVENTOR(S): Black, Michael Terence, Chester Springs, PA, United States  
 Fedon, Jason Craig, Strafford, PA, United States  
 Hodgson, John Edward, Malvern, PA, United States  
 Knowles, David Justin Charles, Boroughbridge, United Kingdom  
 Lonetto, Michael Arthur, Collegeville, PA, United States  
 Kosmatka, Anna Lisa, Doylestown, PA, United States  
 Nicholas, Richard Oakley, Collegeville, PA, United States  
 Palmer, Leslie Marie, Audubon, PA, United States  
 Shilling, Lisa Kathleen, Newtown, PA, United States  
 Stodola, Robert King, Flourtown, PA, United States  
 Warren, Richard Lloyd, Blue Bell, PA, United States  
 PATENT ASSIGNEE(S): SmithKline Beecham Corporation, Philadelphia, PA, United States (U.S. corporation)  
 SmithKline Beecham p.l.c., United Kingdom (non-U.S. corporation)

NUMBER	KIND	DATE
US 5932701		19990803
US 1997-978458		19971125 (8)

PATENT INFORMATION: US 5932701  
 APPLICATION INFO.: US 1997-978458  
 RELATED APPLN. INFO.: Continuation of Ser. No. WO 1997-US14436, filed on 15 Aug 1997 which is a continuation of Ser. No. US 1997-911503, filed on 15 Aug 1997

NUMBER	DATE
US 1996-24022	19960816 (60)

PRIORITY INFORMATION: US 1996-24022  
 DOCUMENT TYPE: Utility  
 FILE SEGMENT: Granted  
 PRIMARY EXAMINER: Minnifield, Nita  
 LEGAL REPRESENTATIVE: King, William T., Gimmi, Edward R., Jackson, Arthur E.  
 NUMBER OF CLAIMS: 1  
 EXEMPLARY CLAIM: 1  
 LINE COUNT: 1726  
 CAS INDEXING IS AVAILABLE FOR THIS PATENT.  
 AB The invention provides riba polypeptides and polynucleotides encoding riba polypeptides and methods for producing such polypeptides by recombinant techniques. Also provided are methods for utilizing riba polypeptides to screen for antibacterial compounds.

09/690,353

L6 ANSWER 21 OF 68 USPATFULL  
 ACCESSION NUMBER: 1999:40236 USPATFULL  
 TITLE: Cell cultures of and cell culturing method for nontransformed pancreatic, thyroid, and parathyroid cells  
 INVENTOR(S): Coon, Hayden G., Gaithersburg, MD, United States  
 Ambesi-Impionbato, Francesco Saverio, Tricesimo, Italy  
 Curcio, Francesco, Pagnacco, Italy  
 PATENT ASSIGNEE(S): Human Cell Cultures Inc., East Sebago, ME, United States (U.S. corporation)

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 5888816		19990330
APPLICATION INFO.:	US 1995-480022		19950607 (8)
RELATED APPLN. INFO.:	Continuation of Ser. No. US 1993-83772, filed on 30 Jun 1993, now abandoned which is a continuation-in-part of Ser. No. US 1993-44010, filed on 8 Apr 1993, now abandoned		
DOCUMENT TYPE:	Utility		
FILE SEGMENT:	Granted		
PRIMARY EXAMINER:	Lankford, Jr., Leon B.		
ASSISTANT EXAMINER:	Tate, Christopher R.		
LEGAL REPRESENTATIVE:	Bundock, John P.		
NUMBER OF CLAIMS:	34		
EXEMPLARY CLAIM:	1		
NUMBER OF DRAWINGS:	18 Drawing Figure(s); 11 Drawing Page(s)		
LINE COUNT:	1992		

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The present invention provides a method for producing an expanded, enriched, non-transformed human cell culture of human pancreatic, thyroid or parathyroid endocrine cells and other types of cells which comprises (1) preparing partially purified, minced tissue that includes a desired type of cells; (2) concentrating the desired cells; (3) resuspending the concentrated cells in a growth medium which selects in favor of the desired cells and in which those cells are proliferated without being transformed and differentiated functions are retained through periodic passaging; (4) culturing the resuspended cells in the growth medium to effect sustained cell division; and (5) passaging the cultured cells periodically to expand the culture. The present invention further provides clonal strains of cells derived from the above-mentioned cell culture and procedures to form matrix-embedded aggregated and non-aggregated cells for providing pseudotissues and products such as matrix-embedded pancreatic islets (pseudoislets). Growth medium and conditioned medium is provided for the culturing of the cells and clonal strains, the growth medium comprising a suitable basal medium supplemented with effective concentrations of hypothalamus and pituitary extracts, serum and other ingredients, which growth medium selects in favor of desired human cells and against passenger cells including fibroblast, macrophage, and capillary endothelial cells such that the desired cells are selectively proliferated without being transformed and an expanded cell culture is provided of functionally differentiated, expanded, non-transformed human cells that is substantially free of such passenger cells.

L6 ANSWER 22 OF 68 USPATFULL  
 ACCESSION NUMBER: 1999:36962 USPATFULL  
 TITLE: Methods for the detection of nitric oxide in fluid media  
 INVENTOR(S): Lai, Ching-San, Encinitas, CA, United States  
 PATENT ASSIGNEE(S): Medinor, Inc., San Diego, CA, United States (U.S. corporation)

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 5885842		19990323
APPLICATION INFO.:	US 1995-745678		19961108 (8)
DOCUMENT TYPE:	Utility		
FILE SEGMENT:	Granted		
PRIMARY EXAMINER:	Green, Lora M.		
ASSISTANT EXAMINER:	Ricigliano, Joseph W.		
LEGAL REPRESENTATIVE:	Gray Cary Ware & Freidenrich, Reiter, Stephen E.		
NUMBER OF CLAIMS:	30		
EXEMPLARY CLAIM:	1		
NUMBER OF DRAWINGS:	4 Drawing Figure(s); 2 Drawing Page(s)		
LINE COUNT:	907		

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB Non-invasive methods have been developed for the measurement of NO levels in a variety of fluid media, e.g., body fluids. The present invention embraces the use of a semi-permeable vessel wherein said vessel contains an NO reacting substance to trap NO diffusing thereinto, and a simple physical or chemical detection method to measure the levels of the end products. Since NO is a neutral gas molecule, it is capable of diffusing freely across a wide range of biocompatible polymer membranes which exhibit a high permeability to NO and other neutral gas molecules, such as O.sub.2 and CO.sub.2, but which are not permeable to charged molecules, such as NO.sub.3.sup.- or NO.sub.2.sup.-. The latter two compounds are ubiquitously present in body fluids and often interfere with the measurement of NO levels. The permeability of selected membranes to NO, but not to NO.sub.3.sup.- or NO.sub.2.sup.-, makes it possible for the bags employed in the practice of the invention to selectively collect NO, even in the presence of potentially competing species. The simple, easy and non-invasive methods of the invention will find a variety of uses, e.g., for diagnosis and monitoring of NO overproduction (and underproduction) that has been associated with many inflammatory and infectious diseases.

L6 ANSWER 21 OF 68 USPATFULL (Continued)

L6 ANSWER 23 OF 68 USPATFULL  
 ACCESSION NUMBER: 1999:19129 USPATFULL  
 TITLE: Methods of receptor modulation and uses therefor  
 INVENTOR(S): Morgan, Jr., A. Charles, Edmonds, WA, United States  
 Wilbur, D. Scott, Edmonds, WA, United States  
 PATENT ASSIGNEE(S): Receptagen Corporation, Edmonds, WA, United States (U.S. corporation)  
 University of Washington, Seattle, WA, United States (U.S. corporation)

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 5869465		19990209
APPLICATION INFO.:	US 1995-406194		19950316 (8)
RELATED APPLN. INFO.:	Continuation-in-part of Ser. No. US 1994-224831, filed on 8 Apr 1994, now abandoned		
DOCUMENT TYPE:	Utility		
FILE SEGMENT:	Granted		
PRIMARY EXAMINER:	Tsang, Cecilia J.		
ASSISTANT EXAMINER:	Gupta, Anish		
LEGAL REPRESENTATIVE:	Christensen O'Connor Johnson & Kindness PLLC		
NUMBER OF CLAIMS:	13		
EXEMPLARY CLAIM:	1		
NUMBER OF DRAWINGS:	28 Drawing Figure(s); 18 Drawing Page(s)		
LINE COUNT:	2882		

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB Receptor modulating agents capable of modulating cell surface receptors by affecting the cell surface receptor trafficking pathway are utilized for the treatment and diagnosis of a variety of disorders in warm-blooded animals, including neoplastic disorders. The receptor modulating agents are comprised of a covalently bound rerouting moiety and targeting moiety.

09/690,353

L6 ANSWER 24 OF 68 USPATFULL  
 ACCESSION NUMBER: 1999:4383 USPATFULL  
 TITLE: PCR A  
 INVENTOR(S): Black, Michael Terence, Chester Springs, PA, United States  
 Hodgeson, John Edward, Malvern, PA, United States  
 Knowles, David Justin Charles, Redhill, United Kingdom  
 Lonetto, Michael Arthur, Collegeville, PA, United States  
 O Nicholas, Richard, Collegeville, PA, United States  
 Stodola, Robert King, Flourtown, PA, United States  
 Holmes, David J., West Chester, PA, United States  
 PATENT ASSIGNEE(S): SmithKline Beecham Corporation, Philadelphia, PA, United States (U.S. corporation)

NUMBER	KIND	DATE
PATENT INFORMATION:	US 5858718	19990112
APPLICATION INFO.:	US 1997-889711	19970708 (8)
DOCUMENT TYPE:	Utility	
FILE SEGMENT:	Granted	
PRIMARY EXAMINER:	Duffy, Patricia	
LEGAL REPRESENTATIVE:	Gimmis, Edward R., King, William T., Jackson, Arthur E.	
NUMBER OF CLAIMS:	11	
EXEMPLARY CLAIM:	1	
LINE COUNT:	1352	

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The invention provides PCR polypeptides and DNA (RNA) encoding PCR polypeptides and methods for producing such polypeptides by recombinant techniques. Also provided are methods for utilizing PCR polypeptides to screen for antibacterial compounds.

L6 ANSWER 25 OF 68 CAPLUS COPYRIGHT 2001 ACS DUPLICATE 2  
 ACCESSION NUMBER: 1998:207280 CAPLUS  
 DOCUMENT NUMBER: 128:275101  
 TITLE: Gas and gaseous precursor filled microspheres as topical and subcutaneous delivery vehicles  
 Unger, Evan C.; Matsunaga, Terry O.; Yellowhair, David  
 INVENTOR(S): Iamarx Pharmaceutical Corp., USA  
 PATENT ASSIGNEE(S): U.S., 40 pp. Cont.-in-part of U.S. Ser. No. 307,305.  
 SOURCE: CODEN: USXXAM  
 DOCUMENT TYPE: Patent  
 LANGUAGE: English  
 FAMILY ACC. NUM. COUNT: 19  
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 5733572	A	19980331	US 1994-346426	19941129
US 5088499	A	19920218	US 1990-569828	19900820
WO 9109629	A1	19910711	WO 1990-US7500	19901219
W: CA, JP				
RW: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LU, NL, SE				
JP 05502675	T2	19930513	JP 1991-503276	19901219
AT 180170	E	19990615	AT 1991-902857	19901219
ES 211051	T3	19990716	ES 1991-902857	19901219
US 5228446	A	19930720	US 1991-717084	19910618
WO 9222247	A1	19921223	WO 1992-US2615	19920331
W: AU, CA, JP				
RW: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LU, MC, NL, SE				
AU 9220020	A1	19930112	AU 1992-20020	19920331
AU 667471	B2	19960328		
JP 06508364	T2	19940922	JP 1992-500847	19920331
EP 616508	A1	19940928	EP 1992-912456	19920331
EP 616508	B1	20010718		
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, MC, NL, SE				
AT 203148	E	20010815	AT 1992-912456	19920331
US 5469854	A	19951128	US 1993-76239	19930611
US 5580575	A	19961203	US 1993-76250	19930611
US 5348016	A	19940920	US 1993-88268	19930707
US 5542935	A	19960806	US 1993-160232	19931130
US 5585112	A	19961217	US 1993-159687	19931130
US 5769080	A	19980623	US 1994-199462	19940222
WO 9428874	A1	19941222	WO 1994-US5633	19940519
W: AU, CA, CN, JP				
RW: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE				
US 5773024	A	19980630	US 1994-307305	19940916
CA 2177713	A	19950608	CA 1994-2177713	19941130
JP 09506098	T2	19970617	JP 1994-515763	19941130
US 5571497	A	19961105	US 1995-468056	19950606
CN 1180310	A	19980429	CN 1996-193069	19960327
US 6001335	A	19991214	US 1996-665719	19960618
US 5935553	A	19990810	US 1996-758179	19961125
US 5985246	A	19991116	US 1997-888426	19970708
AU 713127	B2	19991125	AU 1998-56271	19980224
AU 9856271	A1	19980507		
AU 9888405	A1	19981203	AU 1998-88405	19981012
AU 731072	B2	20010322		

L6 ANSWER 25 OF 68 CAPLUS COPYRIGHT 2001 ACS DUPLICATE 2

(Continued)

AU 9910043 A1 19990304 AU 1999-10043 19990104  
 PRIORITY APPLN. INFO.:  
 US 1989-455707 B2 19891222  
 US 1990-569828 A2 19900820  
 US 1991-716899 B2 19910618  
 US 1991-717084 A2 19910618  
 US 1992-76239 A2 19930611  
 US 1993-76250 A2 19930611  
 US 1993-159674 B2 19931130  
 US 1993-159687 A2 19931130  
 US 1993-160232 A2 19931130  
 US 1994-307305 A2 19940916  
 WO 1990-US7500 W 19901219  
 US 1991-750877 A3 19910826  
 US 1992-818069 A3 19920108  
 WO 1992-US2615 A 19920331  
 US 1992-967974 A3 19921027  
 US 1993-17683 A3 19930212  
 US 1993-18112 B3 19930217  
 US 1993-85608 A3 19930630  
 US 1993-88268 A3 19930707  
 US 1993-309305 A3 19931130  
 US 1993-163039 A3 19931206  
 US 1994-212553 B2 19940311  
 AU 1994-70416 A3 19940519  
 US 1994-346426 19941129  
 AU 1995-21850 A3 19941130  
 WO 1994-US13817 W 19941130  
 US 1995-395683 A3 19950228  
 US 1995-468056 A3 19950606  
 US 1995-471250 A3 19950606  
 US 1996-665719 A3 19960618

AB Gas and gaseous precursor filled microspheres, and foams provide novel topical and s.c. delivery vehicles for various active ingredients, including drugs and cosmetics. Gas and gaseous precursor filled microcapsules were prep'd. from dipalmitoylphosphatidylcholine.

L6 ANSWER 26 OF 68 CAPLUS COPYRIGHT 2001 ACS  
 ACCESSION NUMBER: 1998:55494 CAPLUS  
 DOCUMENT NUMBER: 128:112664  
 TITLE: Pressure-mediated binding of biomolecular complexes  
 Litt, Gerald J.; Laugharn, James A., Jr.; Green, David  
 INVENTOR(S): J.; Hess, Robert A.; Paulus, Henry  
 PATENT ASSIGNEE(S): Bioseq, Inc., USA; Litt, Gerald J.; Laugharn, James A., Jr.; Green, David J.; Hess, Robert A.; Paulus, Henry  
 SOURCE: PCT Int. Appl., 89 pp.  
 CODEN: PIXXD2  
 DOCUMENT TYPE: Patent  
 LANGUAGE: English  
 FAMILY ACC. NUM. COUNT: 1  
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 9800032	A1	19980108	WO 1997-US11198	19970701
W: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GR, HU, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, TJ, TM, TR, TT, UA, UG, US, UZ, VN, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
RW: GH, KE, LE, MW, SD, SE, ZW, AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG				
CA 2259318	AA	19980108	CA 1997-2259318	19970701
AU 9736437	A1	19980121	AU 1997-36437	19970701
EP 924991	A1	19990630	EP 1997-933189	19970701
EP 924991	B1	20011004		
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, FI				

PRIORITY APPLN. INFO.:  
 US 1996-20562 P 19960702  
 US 1997-44595 P 19970422  
 WO 1997-US11198 W 19970701

AB The invention relates to (1) pressure-mediated dissociation of an analyte complexed with an endogenous binding partner to enable detection of a complex formed from the analyte and an exogenous binding factor, (2) pressure-mediated association of an analyte and an exogenous binding partner to enable more rapid and/or more sensitive detection of an analyte, and (3) pressure-mediated association and dissociation of biomol. complexes to enable sepn. of one biomol. from a complex mixt. Pressure can be used to improve assays by dissociating endogenous analyte complexes and improving assay speed and sensitivity by associating the analyte mol. with exogenously supplied binding partners. Pressure can also be used to improve the sepn. of compds. from contaminated mixts.

09/690,353

L6 ANSWER 27 OF 68 USPATFULL  
 ACCESSION NUMBER: 1998:157191 USPATFULL  
 TITLE: Cell cultures of and cells culturing method for nontransformed parotid cells  
 INVENTOR(S): Coon, Hayden G., Gaithersburg, MD, United States  
 Ambesi-Impionbato, Francesco Saverio, Tricesimo, Italy  
 Curcio, Francesco, Pagnacco, Italy  
 PATENT ASSIGNEE(S): Human Cell Cultures Inc., East Sebago, ME, United States (U.S. corporation)

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 5849584		19981215
APPLICATION INFO.:	US 1995-485650		19950607 (8)
RELATED APPLN. INFO.:	Continuation of Ser. No. US 1993-83772, filed on 30 Jun 1993, now abandoned which is a continuation-in-part of Ser. No. US 1993-44010, filed on 8 Apr 1993, now abandoned		

DOCUMENT TYPE: Utility  
 FILE SEGMENT: Granted  
 PRIMARY EXAMINER: Lankford, Jr., Leon B.  
 ASSISTANT EXAMINER: Tate, Christopher R.  
 LEGAL REPRESENTATIVE: Bundock, John P.  
 NUMBER OF CLAIMS: 17  
 EXEMPLARY CLAIM: 1  
 NUMBER OF DRAWINGS: 18 Drawing Figure(s); 11 Drawing Page(s)  
 LINE COUNT: 1832  
 CAS INDEXING IS AVAILABLE FOR THIS PATENT.  
 AB The present invention provides a method for producing an expanded non-transformed cell culture comprising the steps of: (1) preparing partially purified, minced tissue; (2) concentrating the resulting cells and tissue pieces; (3) resuspending the concentrated tissue cells and pieces in a culture medium capable of supporting sustained cell division that is contained in a culture vessel; (4) incubating the cells; and (5) passaging the cells periodically. The present invention further provides clonal strains of cells derived from the above-mentioned cell culture, medium and conditioned medium designed for the culturing of parotid cells and other glandular cells such as pancreatic, thyroid, and parathyroid, and cells, and the use of cultured pancreatic cells to form pancreatic pseudotissues composed of matrix-embedded aggregated (pseudoislets) or individual cells, to treat blood sugar disorders in mammals, and to test for cytotoxicity and autoimmune activities with reference to pancreatic endocrine cells. The nontransformed cells are cultured in a growth medium comprising a suitable basal medium supplemented with effective concentrations of hypothalamus and pituitary extracts, and serum.

L6 ANSWER 29 OF 68 USPATFULL  
 ACCESSION NUMBER: 1998:147427 USPATFULL  
 TITLE: Water soluble vitamin B.sub.12 receptor modulating agents and methods related thereto  
 INVENTOR(S): Morgan, Jr., A. Charles, Mill Creek, WA, United States  
 Wilbur, D. Scott, Edmonds, WA, United States  
 Pathare, Pradip M., Seattle, WA, United States  
 PATENT ASSIGNEE(S): Receptagen Corporation, Edmonds, WA, United States (U.S. corporation)  
 University of WA, Edmonds, WA, United States (U.S. corporation)

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 5840712		19981124
APPLICATION INFO.:	US 1995-545151		19951019 (8)
RELATED APPLN. INFO.:	Continuation-in-part of Ser. No. US 1995-406191, filed on 16 Mar 1995 Ser. No. Ser. No. US 1995-406192, filed on 16 Mar 1995, now patented, Pat. No. US 5739287 And Ser. No. US 1995-406194, filed on 16 Mar 1995, each Ser. No. US which is a continuation-in-part of Ser. No. US 1993-44010, filed on 8 Apr 1993, now abandoned		

No. US 1994-224831, filed on 8 Apr 1994, now abandoned  
 DOCUMENT TYPE: Utility  
 FILE SEGMENT: Granted  
 PRIMARY EXAMINER: Mutzell, Paula K.  
 ASSISTANT EXAMINER: Bakalyar, Heather A.  
 LEGAL REPRESENTATIVE: Christensen O'Connor Johnson & Kindness PLLC  
 NUMBER OF CLAIMS: 10  
 EXEMPLARY CLAIM: 1  
 NUMBER OF DRAWINGS: 28 Drawing Figure(s); 18 Drawing Page(s)  
 LINE COUNT: 3615  
 CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB Vitamin B.sub.12 receptor modulating agents capable of modulating cell surface receptors by affecting the cell surface receptor trafficking pathway are disclosed. The vitamin B.sub.12 receptor modulating agents are comprised of a covalently bound rerouting moiety and targeting moiety linked by a water-solubilizing linker.

L6 ANSWER 28 OF 68 USPATFULL  
 ACCESSION NUMBER: 1998:147590 USPATFULL  
 TITLE: Receptor modulating agents  
 INVENTOR(S): Morgan, Jr., A. Charles, Edmonds, WA, United States  
 Wilbur, D. Scott, Edmonds, WA, United States  
 PATENT ASSIGNEE(S): Receptagen Corporation, Edmonds, WA, United States (U.S. corporation)  
 University of Washington, Seattle, WA, United States (U.S. corporation)

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 5840880		19981124
APPLICATION INFO.:	US 1995-406191		19950316 (8)
RELATED APPLN. INFO.:	Continuation-in-part of Ser. No. US 1994-224831, filed on 8 Apr 1994, now abandoned		

DOCUMENT TYPE: Utility  
 FILE SEGMENT: Granted  
 PRIMARY EXAMINER: Robinson, Douglas  
 ASSISTANT EXAMINER: Gupta, Anish  
 LEGAL REPRESENTATIVE: Christensen O'Connor Johnson & Kindness PLLC  
 NUMBER OF CLAIMS: 13  
 EXEMPLARY CLAIM: 1  
 NUMBER OF DRAWINGS: 26 Drawing Figure(s); 20 Drawing Page(s)  
 LINE COUNT: 2940  
 CAS INDEXING IS AVAILABLE FOR THIS PATENT.  
 AB Receptor modulating agents capable of modulating cell surface receptors by affecting the cell surface receptor trafficking pathway. The receptor modulating agents are comprised of a covalently bound rerouting moiety and targeting moiety.

L6 ANSWER 30 OF 68 USPATFULL  
 ACCESSION NUMBER: 1998:82596 USPATFULL  
 TITLE: Method of altering blood sugar levels using non-transformed human pancreatic cells that have been expanded in culture  
 INVENTOR(S): Coon, Hayden G., Gaithersburg, MD, United States  
 Ambesi-Impionbato, Francesco Saverio, Tricesimo, Italy  
 Curcio, Francesco, Pagnacco, Italy  
 PATENT ASSIGNEE(S): Human Cell Cultures Inc., East Sebago, ME, United States (U.S. corporation)

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 5780299		19980714
APPLICATION INFO.:	US 1995-480027		19950607 (8)
RELATED APPLN. INFO.:	Division of Ser. No. US 1993-83772, filed on 30 Jun 1993, now abandoned which is a continuation-in-part of Ser. No. US 1993-44010, filed on 8 Apr 1993, now abandoned		

DOCUMENT TYPE: Utility  
 FILE SEGMENT: Granted  
 PRIMARY EXAMINER: Lankford, Jr., Leon B.  
 ASSISTANT EXAMINER: Tate, Christopher R.  
 LEGAL REPRESENTATIVE: Bundock, John P.  
 NUMBER OF CLAIMS: 14  
 EXEMPLARY CLAIM: 1  
 NUMBER OF DRAWINGS: 18 Drawing Figure(s); 11 Drawing Page(s)  
 LINE COUNT: 1828  
 CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The present invention provides a method for producing an expanded non-transformed cell culture comprising the steps of: (1) preparing partially purified, minced tissue; (2) concentrating the resulting cells and tissue pieces; (3) resuspending the concentrated tissue cells and pieces in a culture medium capable of supporting sustained cell division that is contained in a culture vessel; (4) incubating the cells; and (5) passaging the cells periodically. The present invention further provides clonal strains of cells derived from the above-mentioned cell culture, medium and conditioned medium designed for the culturing of such cells, including pancreatic, thyroid, parathyroid, and parotid cells, and the use of cultured pancreatic cells to form pancreatic pseudotissues composed of matrix-embedded aggregated (pseudoislets) or individual cells, to treat blood sugar disorders in mammals, and to test for cytotoxicity and autoimmune activities with reference to pancreatic endocrine cells.

09/690,353

L6 ANSWER 31 OF 68 USPATFULL  
 ACCESSION NUMBER: 1998:39675 USPATFULL  
 TITLE: Biotinylated cobalamins  
 INVENTOR(S): Wilbur, D. Scott, Edmonds, WA, United States  
 Pathare, Pradip M., Seattle, WA, United States  
 Morgan, Jr., A. Charles, Camino Island, WA, United States  
 PATENT ASSIGNEE(S): University of Washington, Seattle, WA, United States (U.S. corporation)  
 Receptagen Corp., Edmonds, WA, United States (U.S. corporation)

NUMBER	KIND	DATE
US 5739287		19980414
US 1995-406192		19950316 (8)

RELATED APPLN. INFO.: Continuation-in-part of Ser. No. US 1994-224831, filed on 8 Apr 1994, now abandoned

DOCUMENT TYPE: Utility  
 FILE SEGMENT: Granted  
 PRIMARY EXAMINER: Russel, Jeffrey E.  
 LEGAL REPRESENTATIVE: Christensen O'Connor Johnson & Kindness PLLC  
 NUMBER OF CLAIMS: 5  
 EXEMPLARY CLAIM: 1  
 NUMBER OF DRAWINGS: 28 Drawing Figure(s); 18 Drawing Page(s)  
 LINE COUNT: 3099

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB A biotinylated cobalamin, formed from a vitamin B.sub.12 molecule coupled to a biotin molecule, is disclosed. In a preferred embodiment, the vitamin B.sub.12 molecule is cyanocobalamin. The biotin molecule can also be coupled to a rerouting moiety, optionally through a biotin binding protein such as avidin or streptavidin. The biotinylated cobalamin binds to a cell surface receptor, is internalized, and once internalized affects the receptor trafficking pathway.

L6 ANSWER 33 OF 68 USPATFULL  
 ACCESSION NUMBER: 97:106797 USPATFULL  
 TITLE: Anti-receptor and growth blocking antibodies to the Vitamin B.sub.12 /transcobalamin II receptor and binding sites  
 INVENTOR(S): Morgan, Jr., Alton Charles, Edmonds, WA, United States  
 PATENT ASSIGNEE(S): Receptagen Corporation, Edmonds, WA, United States (U.S. corporation)

NUMBER	KIND	DATE
US 5688504		19971118
US 1994-306504		19940913 (8)

RELATED APPLN. INFO.: Continuation-in-part of Ser. No. US 1992-880540, filed on 8 May 1992, now abandoned

DOCUMENT TYPE: Utility  
 FILE SEGMENT: Granted  
 PRIMARY EXAMINER: Nucker, Christine M.  
 ASSISTANT EXAMINER: Stucker, Jeffrey  
 LEGAL REPRESENTATIVE: Seed and Berry LLP  
 NUMBER OF CLAIMS: 21  
 EXEMPLARY CLAIM: 1  
 NUMBER OF DRAWINGS: 8 Drawing Figure(s); 8 Drawing Page(s)  
 LINE COUNT: 1860

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB There is disclosed anti-receptor and growth blocking agents to the vitamin B.sub.12 /transcobalamin II receptor and binding sites. The anti-receptor and growth blocking agents antagonize or modulate the vitamin B.sub.12 /transcobalamin II receptor or binding sites, causing cellular depletion of vitamin B.sub.12, thus preventing or inhibiting cell division or causing apoptosis. Anti-receptor and growth blocking agents of the present invention include proteins (such as antibodies and antibody derivatives), peptides and small organic molecules. In a preferred embodiment, the anti-receptor agent is an antibody to the vitamin B.sub.12 /transcobalamin II receptor.

L6 ANSWER 32 OF 68 EMBASE COPYRIGHT 2001 ELSEVIER SCI. B.V.  
 ACCESSION NUMBER: 1998341676 EMBASE  
 TITLE: Transmethylation reactions and autoradiographic distribution of vitamin B12: Effects of clioquinol treatment in mice.  
 AUTHOR: Yassin M.S.; Ekblom J.; Lofberg C.; Orland L.  
 CORPORATE SOURCE: L. Orland, Dept. of Neuroscience (Pharmacology), Biomedical Center, Uppsala University, Box 593, S-751 24 Uppsala, Sweden  
 SOURCE: Japanese Journal of Pharmacology, (1998) 78/1 (55-61).  
 Refs: 23  
 ISSN: 0021-5198 CODEN: JPPAAZ  
 COUNTRY: Japan  
 DOCUMENT TYPE: Journal; Article  
 FILE SEGMENT: 030 Pharmacology  
 037 Drug Literature Index  
 052 Toxicology  
 LANGUAGE: English  
 SUMMARY LANGUAGE: English

AB The catastrophic epidemic of subacute myelo-optic neuropathy (SMON) affected Japan around 1970 with thousands of victims. The cause was attributed to high doses of locally acting oxyquinolines. It has been speculated that oxyquinoline derivatives of the clioquinol type can disturb the retention of vitamin B12 through chelation of Co2+. In the present paper, possible effects of clioquinol on the uptake and tissue distribution of [57Co]-cyanocobalamin have been studied in mice.

In vivo experiments showed markedly decreased accumulation of radiolabelled vitamin B12 in the kidney and skin in animals that were pre-treated with clioquinol. The chloroform:water partition coefficients for [57Co]-cyanocobalamin in the presence or absence of clioquinol were also determined. No statistically significant alterations in the partition coefficient for [57Co]-cyanocobalamin in the presence of clioquinol was evident, indicating that clioquinol does not bind cobalt. In addition, transmethylation reactions in the CNS in mice treated with clioquinol were studied. Specific activities of methionine adenosyltransferase, and S-adenosylhomocysteine levels were not affected. In contrast, clioquinol treatment caused a significant increase in the levels of S-adenosylmethionine in the brain. The data of the present study show that clioquinol treatment can affect the accumulation of vitamin B12 in the kidney and the skin but not in the brain. These results do not support the hypothesis that clioquinol causes its damage to the nervous system by a direct chemical interaction with vitamin B12.

L6 ANSWER 34 OF 68 USPATFULL  
 ACCESSION NUMBER: 97:91199 USPATFULL  
 TITLE: Nanoparticles containing an active substance and a ketalized polytartramic acid, process for their preparation, and use thereof  
 INVENTOR(S): Ahlers, Michael, Mainz, Germany, Federal Republic of  
 Walch, Axel, Frankfurt am Main, Germany, Federal Republic of  
 Seipke, Gerhard, Hofheim, Germany, Federal Republic of  
 Russell-Jones, Gregory, Middel Cove, Australia  
 PATENT ASSIGNEE(S): Hoechst Aktiengesellschaft, Frankfurt am Main, Germany, Federal Republic of (non-U.S. corporation)

NUMBER	KIND	DATE
US 5674531		19971007
US 1995-399474		19950307 (8)

PATENT INFORMATION: DE 1994-4407898 19940309  
 DOCUMENT TYPE: Utility  
 FILE SEGMENT: Granted  
 PRIMARY EXAMINER: Page, Thurman K.  
 ASSISTANT EXAMINER: Spear, James M.  
 LEGAL REPRESENTATIVE: Finnegan, Henderson, Farabow, Garrett & Dunner, L.L.P.  
 NUMBER OF CLAIMS: 11  
 EXEMPLARY CLAIM: 1  
 LINE COUNT: 571

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB Nanoparticles containing an active substance and a ketalized polytartramic acid, process for their preparation, and use thereof. Nanoparticles containing an active substance and a ketalized polytartramic acid are suitable as vehicles for active substances, in particular for peptides and proteins. Processes for the preparation of the nanoparticles are described.

09/690,353

L6 ANSWER 35 OF 68 USPATFULL  
 ACCESSION NUMBER: 97:83944 USPATFULL  
 TITLE: Methods of treating neurological diseases and etiologically related symptomology using carbonyl trapping agents in combination with previously known medicaments  
 INVENTOR(S): Shapiro, Howard K., 214 Price Ave. F32, Narberth, PA, United States 19072

NUMBER	KIND	DATE
US 5668117		19970916
US 1993-62201		19930629 (8)

RELATED APPLN. INFO.: Continuation-in-part of Ser. No. US 1993-26617, filed on 23 Feb 1993, now abandoned which is a continuation of Ser. No. US 1991-660561, filed on 22 Feb 1991, now abandoned

DOCUMENT TYPE: Utility  
 FILE SEGMENT: Granted  
 PRIMARY EXAMINER: Kight, John  
 ASSISTANT EXAMINER: Leary, Louise  
 LEGAL REPRESENTATIVE: Perrella, D. J.  
 NUMBER OF CLAIMS: 29  
 EXEMPLARY CLAIM: 1  
 LINE COUNT: 3963

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB Therapeutic compositions comprising an effective amount of at least one carbonyl trapping agent alone or in combination with a therapeutically effective of a co-agent or medicament are disclosed. The compositions are used to treat a mammal suffering from a neurological disease characterized by covalent bond crosslinking between the nerve cells, other cellular structures and their intracellular and extracellular components, with disease induced carbonyl-containing aliphatic or aromatic hydrocarbons present in mammals.

L6 ANSWER 36 OF 68 USPATFULL  
 ACCESSION NUMBER: 97:78345 USPATFULL  
 TITLE: Nucleophilic polysubstituted aryl acridinium ester conjugates and syntheses thereof  
 INVENTOR(S): Law, Say-Jong, Westwood, MA, United States  
 PATENT ASSIGNEE(S): Chiron Diagnostics Corporation, Malpole, MA, United States (U.S. corporation)

NUMBER	KIND	DATE
US 5663074		19970902
US 1993-32947		19930317 (8)

RELATED APPLN. INFO.: Continuation-in-part of Ser. No. US 1992-871601, filed on 17 Apr 1992 which is a continuation of Ser. No. US 1988-249520, filed on 26 Sep 1988, now abandoned

DOCUMENT TYPE: Utility  
 FILE SEGMENT: Granted  
 PRIMARY EXAMINER: Woodward, Michael P.  
 LEGAL REPRESENTATIVE: Morgenstern, Arthur S., Blackburn, Robert P., Klee, Maurice M.  
 NUMBER OF CLAIMS: 53  
 EXEMPLARY CLAIM: 1  
 NUMBER OF DRAWINGS: 18 Drawing Figure(s); 15 Drawing Page(s)  
 LINE COUNT: 1554

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB This invention is directed to novel nucleophilic polysubstituted aryl acridinium conjugates and the methods for preparation thereof. The nucleophilic polysubstituted aryl acridinium conjugates are useful in biological assays, including novel assays for the determination of Vitamin B.sub.12, folate, cortisol, estradiol, and thromboxane B.sub.2.

L6 ANSWER 37 OF 68 USPATFULL  
 ACCESSION NUMBER: 97:59097 USPATFULL  
 TITLE: Method for preparing an expanded culture and clonal strains of pancreatic, thyroid or parathyroid cells  
 INVENTOR(S): Coon, Hayden G., Gaithersburg, MD, United States  
 Ambei-Impionato, Francesco Saverio, Tricesimo, Italy  
 Curcio, Francesco, Pagnacco, Italy  
 PATENT ASSIGNEE(S): Human Cell Cultures, Inc., Gaithersburg, MD, United States (U.S. corporation)

NUMBER	KIND	DATE
US 5646035		19970708
US 1995-480149		19950607 (8)

RELATED APPLN. INFO.: Continuation of Ser. No. US 1993-83772, filed on 30 Jun 1993, now abandoned which is a continuation-in-part of Ser. No. US 1993-44010, filed on 8 Apr 1993, now abandoned

DOCUMENT TYPE: Utility  
 FILE SEGMENT: Granted  
 PRIMARY EXAMINER: Rollins, John W.  
 ASSISTANT EXAMINER: Larson, Kristin  
 LEGAL REPRESENTATIVE: Leydig, Voit & Mayer, Ltd.  
 NUMBER OF CLAIMS: 16  
 EXEMPLARY CLAIM: 1  
 NUMBER OF DRAWINGS: 18 Drawing Figure(s); 11 Drawing Page(s)  
 LINE COUNT: 1831

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The present invention provides a method for producing an expanded non-transformed cell culture comprising the steps of: (1) preparing partially purified, minced tissue; (2) concentrating the resulting cells and tissue pieces; (3) resuspending the concentrated tissue cells and pieces in a culture medium capable of supporting sustained cell division that is contained in a culture vessel; (4) incubating the cells; and (5) passaging the cells periodically. The present invention further provides clonal strains of cells derived from the above-mentioned cell culture, medium and conditioned medium designed for the culturing of such cells, including pancreatic, thyroid, parathyroid, and parotid cells, and the use of cultured pancreatic cells to form pancreatic pseudotissues composed of matrix-embedded aggregated (pseudotissues) or individual cells, to treat blood sugar disorders in mammals, and to test for cytotoxicity and autoimmune activities with reference to pancreatic endocrine cells.

L6 ANSWER 38 OF 68 MEDLINE  
 ACCESSION NUMBER: 1998012901 MEDLINE  
 DOCUMENT NUMBER: 98012901 PubMed ID: 9352852  
 TITLE: Apical expression of functional asialoglycoprotein receptor  
 AUTHOR: Mu J Z; Gordon M; Shao J S; Alpers D H  
 CORPORATE SOURCE: Department of Internal Medicine, Washington University School of Medicine, Saint Louis, Missouri, USA.  
 CONTRACT NUMBER: PO1-33487  
 SOURCE: GASTROENTEROLOGY, (1997 Nov) 113 (5) 1501-9.  
 PUB. COUNTRY: United States  
 LANGUAGE: English  
 FILE SEGMENT: Abridged Index Medicus Journals; Priority Journals  
 ENTRY MONTH: 199711  
 ENTRY DATE: Entered STN: 19971224  
 Last Updated on STN: 19971224  
 Entered Medline: 19971113

AB BACKGROUND & AIMS: The asialoglycoprotein receptor localizes to the basolateral membrane of hepatocytes and to the apical membrane of enterocytes. The aim of this study was to examine HT-29 cells as a polarized cell model for studying apically localized endogenous asialoglycoprotein receptor. METHODS: Subunits H1 and H2 (human) were detected by Western blot and immunoprecipitated using subunit-specific antisera against hepatic receptor peptides. Receptor function was assessed by uptake of iodinated asialo-orosomucoid, immunoglobulin (Ig) A1, and haptocorrin. Immunocytochemistry was analyzed by standard light and confocal microscopy. RESULTS: Receptor content of the minor subunit, H2, was predominant. HT-29 cells mediated specific uptake and degradation of 125I-asialo-orosomucoid. A high-affinity (0.6 x 10<sup>-9</sup> mmol/L) and a low-affinity binding site were present. The specific ligand binding capacity of the apical surface was approximately twice that of the basolateral surface. Immunocytochemistry revealed a predominant apical membrane location of the minor receptor subunit, with some intracellular receptor. The apical H2 subunit was preferentially basolaterally located subunit. Human IgA1 bound specifically to HT-29 cells with a molar ratio of 0.26 compared with asialo-orosomucoid; porcine haptocorrin bound with a molar ratio of 1.35. CONCLUSIONS: HT-29 cells produce a functionally apically located asialoglycoprotein receptor and provide a model for receptor trafficking in the enterocyte.

09/690,353

L6 ANSWER 39 OF 68 USPATFULL  
 ACCESSION NUMBER: 96:99222 USPATFULL  
 TITLE: Combination medications containing alpha-lipoic acid and related  
 INVENTOR(S): Weischer, Carl-Heinrich, Bonn, Germany, Federal Republic of  
 Ulrich, Heinz, Niedernberg, Germany, Federal Republic of  
 PATENT ASSIGNEE(S): Wessel, Klaus, Frankfurt, Germany, Federal Republic of  
 Asta Medica Aktiengesellschaft, Dresden, Germany, Federal Republic of (non-U.S. corporation)

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 5569670		19961029
APPLICATION INFO.:	US 1995-404153		19950314 (8)
RELATED APPLN. INFO.:	Division of Ser. No. US 1994-197643, filed on 10 Feb 1994, now abandoned which is a continuation-in-part of Ser. No. US 1993-71259, filed on 4 Jun 1993, now abandoned		

PRIORITY INFORMATION: DE 1992-4218572 19920605  
 DOCUMENT TYPE: Utility  
 FILE SEGMENT: Granted  
 PRIMARY EXAMINER: Dees, Jos e G.  
 ASSISTANT EXAMINER: Lambkin, Deborah  
 LEGAL REPRESENTATIVE: Cushman Darby & Cushman, LLP  
 NUMBER OF CLAIMS: 1  
 EXEMPLARY CLAIM: 1  
 LINE COUNT: 1013  
 CAS INDEXING IS AVAILABLE FOR THIS PATENT.  
 AB A pharmaceutical composition containing alpha-lipoic acid, dihydro-lipoic acid, metabolites of alpha-lipoic acid (inter alia bisnortetralipoic acid and tetranorlipoic acid), optical isomers R- and S- forms of alpha-lipoic acid in oxidized and reduced form together with a vitamin, especially vitamins A, B1, B2, B6, B12, C and E and their pharmaceutically acceptable salts. The compositions are useful for producing analgesic, anti-inflammatory, antidiabetic, cytoprotective, anti-ulcer, antineurotic, neuroprotective, detoxifying, antischemic, liver function regulating, anti-allergic, immune-stimulating and antioncogenic effects.

L6 ANSWER 41 OF 68 USPATFULL  
 ACCESSION NUMBER: 95:64857 USPATFULL  
 TITLE: Folate immunoassay utilizing folate binding protein in a monoclonal antibody format  
 INVENTOR(S): Beggs, Michael J., Waukegan, IL, United States  
 Sohn, Linda J., Palatine, IL, United States  
 Herrmann, Robert J., Gurnee, IL, United States  
 Hau, Stephen, Vernon Hills, IL, United States  
 Hawksworth, David J., Mundelein, IL, United States  
 Pinkus, Mary S., Chicago, IL, United States  
 PATENT ASSIGNEE(S): Abbott Laboratories, Abbott Park, IL, United States (U.S. corporation)

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 5434087		19950718
APPLICATION INFO.:	US 1993-21942		19930224 (8)
DOCUMENT TYPE:	Utility		
FILE SEGMENT:	Granted		
PRIMARY EXAMINER:	Scheiner, Toni R.		
ASSISTANT EXAMINER:	Parsons, Nancy J.		
LEGAL REPRESENTATIVE:	Weinstein, David L.		
NUMBER OF CLAIMS:	25		
EXEMPLARY CLAIM:	1		
NUMBER OF DRAWINGS:	3 Drawing Figure(s); 2 Drawing Page(s)		
LINE COUNT:	711		

CAS INDEXING IS AVAILABLE FOR THIS PATENT.  
 AB An improved method for performing immunoassays whereby specific binding proteins for vitamin B12, folate and other target analytes are utilized with antibodies with different specificities for the binding proteins. Antibodies bridge the specific binding protein directly or indirectly to a capturable material.

L6 ANSWER 40 OF 68 USPATFULL  
 ACCESSION NUMBER: 96:65493 USPATFULL  
 TITLE: Nucleophilic polyubstituted aryl acridinium ester conjugates uses thereof  
 INVENTOR(S): Law, Say-Jong, Westwood, MA, United States  
 Chang, Steve C. S., Franklin, MA, United States  
 Klukas, Carol K., Pittsburgh, PA, United States  
 Vitkauskas, Christine A., North Attleboro, MA, United States  
 PATENT ASSIGNEE(S): Ciba Corning Diagnostics Corp., Medfield, MA, United States (U.S. corporation)

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 5538901		19960723
APPLICATION INFO.:	US 1994-292946		19940818 (8)
RELATED APPLN. INFO.:	Continuation of Ser. No. US 1993-32085, filed on 17 Mar 1993, now abandoned which is a division of Ser. No. US 1992-871601, filed on 17 Apr 1992, now patented, Pat. No. US 5241070, issued on 13 Aug 1993 which is a continuation of Ser. No. US 1988-249620, filed on 26 Sep 1988, now abandoned		

DOCUMENT TYPE: Utility  
 FILE SEGMENT: Granted  
 PRIMARY EXAMINER: Spiegel, Carol A.  
 LEGAL REPRESENTATIVE: Morgenstern, Arthur S., Roesler, Judith A.  
 NUMBER OF CLAIMS: 15  
 EXEMPLARY CLAIM: 1  
 NUMBER OF DRAWINGS: 18 Drawing Figure(s); 15 Drawing Page(s)  
 LINE COUNT: 1444  
 CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB This invention is directed to the novel assay methods utilizing nucleophilic polyubstituted aryl acridinium ester conjugates as the tracers. Conjugates prepared by covalent coupling of novel nucleophilic polyubstituted aryl acridinium esters with biological compounds including small organic molecules such as Vitamin B12, folate, estradiol, and thromboxane B2, were found useful in the development of highly sensitive assays for the analytes of diagnostic interest.

L6 ANSWER 42 OF 68 BIOSIS COPYRIGHT 2001 BIOSIS  
 ACCESSION NUMBER: 1995:509032 BIOSIS  
 DOCUMENT NUMBER: PREV199598514082  
 TITLE: Inhibition of chondrocyte cathepsin B and L activities by insulin-like growth factor-II (IGF-II) and its Ser-29 variant in vitro: Possible role of the mannose 6-phosphate/IGF-II receptor.  
 AUTHOR(S): De Ceuninck, F.; Poiraudou, S.; Pagano, M.; Tsagris, L.; Blanchard, O.; Willeput, J.; Corvol, M. (1)  
 CORPORATE SOURCE: (1) Institut National de la Sante et de la Recherche Medicale, U 30, 149 rue de Sevres, 75743 Paris, Cedex 15 France  
 SOURCE: Molecular and Cellular Endocrinology, (1995) Vol. 113, No. 2, pp. 205-213.  
 ISSN: 0303-7207.  
 DOCUMENT TYPE: Article  
 LANGUAGE: English

AB Lysosomal enzymes and IGF-II both bind to the mannose 6-phosphate (M6P)/IGF-II receptor. This receptor targets newly synthesized lysosomal enzymes to lysosomes. The functional meaning of IGF-II binding to this receptor is not well known. We have postulated that IGF-II may affect the targeting of lysosomal enzymes in cartilage remodeling. We therefore examined the effect of IGF-II, the Ser-29 IGF-II variant (vIGF-II) and IGF-I on lysosomal cathepsin B and L activities from post-natal rabbit chondrocytes in vitro. This effect was compared with the ability of each peptide to stimulate chondrocyte-sulfated proteoglycan synthesis. The sulfating dose-response relationship of the IGF peptides corresponded to their relative binding affinities for the type I-IGF receptor (IGF-I gt IGF-II gt vIGF-II). The intracellular cathepsin B and

L activities were inhibited in a time- and dose-dependent manner by IGF-II or vIGF-II. Maximal inhibition of cathepsin B and L activities (40 and 30% below controls, respectively) was found after an 8 h treatment with 100 ng/ml IGF-II or vIGF-II. By contrast, IGF-I up to 1 mu-g/ml or insulin up to 2 mu-g/ml had no inhibitory effect. The relative potency pattern corresponded to the binding profile of each ligand for the M6P/IGF-II receptor. A treatment of chondrocytes with IGF-I or insulin transiently increased the binding of radiolabeled IGF-II at the cell surface to -120% of controls, whereas IGF-II or vIGF-II had no effect. Thus, it is unlikely that the inhibition of lysosomal enzyme activities by IGF-II peptides could result from a redistribution of M6P/IGF-II receptors from intracellular compartments to the plasma membrane. We hypothesize that internalized IGF-II peptides could occupy the intracellular M6P/IGF-II binding sites required for the targeting of cathepsins B and L to lysosomes.

09/690,353

L6 ANSWER 43 OF 68 CAPLUS COPYRIGHT 2001 ACS  
 ACCESSION NUMBER: 1994:442743 CAPLUS  
 DOCUMENT NUMBER: 121:42743  
 TITLE: Microparticle formulations containing biodegradable mixed polymers  
 INVENTOR(S): Fritzsche, Thomas; Heldmann, Dieter; Weitschies, Werner  
 PATENT ASSIGNEE(S): Schering A.-G., Germany  
 SOURCE: Ger. Offen., 8 pp.  
 CODEN: GWXXBX  
 DOCUMENT TYPE: Patent  
 LANGUAGE: German  
 FAMILY ACC. NUM. COUNT: 1  
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
DE 4232755	A1	19940331	DE 1992-4232755	19920926
WO 9407539	A1	19940414	WO 1993-EP2422	19930908
W: AU, CA, FI, HU, JP, KR, NO, NZ, US				
RW: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE				
EP 662005	A1	19950712	EP 1993-919293	19930908
EP 662005	B1	19980429		
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LI, LU, MC, NL, PT, SE				
HU 71683	A2	19960129	HU 1995-876	19930908
JP 08504398	B2	19960514	JP 1993-508620	19930908
AU 687360	B2	19980226	AU 1993-49592	19930908
AT 165517	E	19980515	AT 1993-919293	19930908
ES 2118251	T3	19980916	ES 1993-919293	19930908
ZA 9307099	A	19950324	ZA 1993-7029	19930924
CN 1089508	A	19940720	CN 1993-118159	19930925
IL 107108	A1	19980104	IL 1993-107108	19930926
FI 9501379	A	19950323	FI 1995-1379	19950323
NO 9501138	A	19950324	NO 1995-1138	19950324
PRIORITY APPLN. INFO.:			DE 1992-4232755	19920926
			WO 1993-EP2422	19930908

AB Microcapsules for diagnostic or therapeutic use are prep'd. comprising (1) a wall of biopolymer/synthetic polymer interpolymer, which has site-specific phys. properties or bears functional groups for binding e.g. chelating ligands or their metal complexes, and (2) an optional core consisting of a gas or therapeutic agent. Thus, an autoclaved gelatin soln. was adjusted to pH 5.0 and stirred with Bu cyanacrylate to produce air-filled dispersible microparticles [55 mol% gelatin, 45 mol% poly(Bu cyanacrylate)] for use in contrast media for sonog.

L6 ANSWER 45 OF 68 MEDLINE  
 ACCESSION NUMBER: 95070174 MEDLINE  
 DOCUMENT NUMBER: 95070174 PubMed ID: 7979409  
 TITLE: Effect of processing inhibitors on cobalamin (vitamin B12) transcytosis in polarized opossum kidney cells.  
 AUTHOR: Ramanujam K S; Seetharam S; Dahma N M; Seetharam B  
 CORPORATE SOURCE: Department of Medicine, Medical College of Wisconsin, Milwaukee 53226.  
 CONTRACT NUMBER: DK-26638 (NIDDK)  
 SOURCE: ARCHIVES OF BIOCHEMISTRY AND BIOPHYSICS, (1994 Nov 15) 315 (1) 8-15.  
 PUB. COUNTRY: Journal code: 6SK; 0372430. ISSN: 0003-9861.  
 LANGUAGE: United States  
 FILE SEGMENT: Journal; Article; (JOURNAL ARTICLE)  
 ENTRY MONTH: English  
 ENTRY DATE: Priority Journals  
 199412  
 Entered STN: 19950110  
 Last Updated on STN: 19950110  
 Entered Medline: 19941216

AB The main objectives of the current study were to investigate the effect of tunicamycin and other posttranslational processing inhibitors on the apical brush border expression of intrinsic factor-cobalamin receptor (IFCR) and the apical to basolateral transcytosis of cobalamin (Cbl). Because of the high and selective expression of IFCR in the apical brush border membrane of opossum kidney (OK) cells (K. S. Ramanujam, S. Seetharam, N. Dahma, and B. Seetharam, (1991) J. Biol. Chem. 266, 13135-13140), we have used cultured OK cells to address these issues. When polarized OK cells grown on culture inserts were incubated with tunicamycin, deoxynojirimycin, swainsonine, or cerulenin, the surface binding of the ligand, intrinsic factor-[56Co]Cbl was inhibited by tunicamycin but not by the other inhibitors. However, Cbl transcytosis was inhibited by both tunicamycin and cerulenin but not with deoxynojirimycin or swainsonine. Incubation of cells with tunicamycin decreased the half-life of IFCR from 48 to 24 h, thus causing faster degradation and depletion of the surface receptor. Incubation of cells with cerulenin resulted in the intralysosomal retention of internalized Cbl. Mature receptor labeled with either [35S]methionine or [3H]mannose was sensitive to digestion with both endoglycosidase H and peptide N-glycosidase F and revealed the presence of two or three N-linked oligosaccharides of the high mannose or hybrid type. Metabolic labeling of OK cells with [3H]palmitic acid revealed that IFCR was palmitoylated and the label was sensitive to treatment with hydroxylamine. Based on these results we suggest that IFCR expression in the apical membrane and Cbl transcytosis in polarized OK cells are regulated by core N-glycosylation but not by further processing of the terminal sugars. In addition, we also suggest that the inhibition of Cbl transcytosis by cerulenin is due to inhibition of postinternalization events.

L6 ANSWER 44 OF 68 USPATFULL  
 ACCESSION NUMBER: 94:88598 USPATFULL  
 TITLE: Lyophilized ligand-receptor complexes for assays and sensors  
 INVENTOR(S): Ligler, Frances S., Potomac, MD, United States  
 Whelan, James P., Potomac, MD, United States  
 PATENT ASSIGNEE(S): The United States of America as Represented by the Secretary of the Navy, Washington, DC, United States (U.S. government)  
 U.S. Drug Testing, Inc., Rancho Cucamonga, CA, United States (U.S. corporation)

NUMBER	KIND	DATE
US 5354654		19941011
US 1993-92518		19930716 (8)

PATENT INFORMATION: US 5354654  
 APPLICATION INFO.: US 1993-92518  
 DOCUMENT TYPE: Utility  
 FILE SEGMENT: Granted  
 PRIMARY EXAMINER: Saunders, David  
 LEGAL REPRESENTATIVE: Oblon, Spivak, McClelland, Maier & Neustadt  
 NUMBER OF CLAIMS: 22  
 EXEMPLARY CLAIM: 1  
 NUMBER OF DRAWINGS: 2 Drawing Figure(s); 2 Drawing Page(s)  
 LINE COUNT: 1129  
 CAS INDEXING IS AVAILABLE FOR THIS PATENT.  
 AB A dry reagent prepared by lyophilizing a labelled ligand-immobilized receptor complex or a labelled receptor-immobilized ligand complex is, on rehydration, useful for detecting analytes in samples in a variety of displacement assays.

L6 ANSWER 46 OF 68 USPATFULL  
 ACCESSION NUMBER: 93:72216 USPATFULL  
 TITLE: Nucleophilic polysubstituted aryl acridinium esters and uses thereof  
 INVENTOR(S): Law, Say-Jong, Westwood, MA, United States  
 PATENT ASSIGNEE(S): Ciba Corning Diagnostic Corp., Medfield, MA, United States (U.S. corporation)

NUMBER	KIND	DATE
US 5241070		19930831
US 1992-871601		19920417 (7)

PATENT INFORMATION: US 5241070  
 APPLICATION INFO.: US 1992-871601  
 DISCLAIMER DATE: 20070417  
 RELATED APPLN. INFO.: Continuation of Ser. No. US 1988-249620, filed on 26 Sep 1988, now abandoned  
 DOCUMENT TYPE: Utility  
 FILE SEGMENT: Granted  
 PRIMARY EXAMINER: Daus, Donald G.  
 LEGAL REPRESENTATIVE: Morgenstern, Arthur S., Slepchuk, Jr., Nicholas I.  
 NUMBER OF CLAIMS: 3  
 EXEMPLARY CLAIM: 1  
 NUMBER OF DRAWINGS: 18 Drawing Figure(s); 15 Drawing Page(s)  
 LINE COUNT: 1023  
 CAS INDEXING IS AVAILABLE FOR THIS PATENT.  
 AB This invention is directed to novel nucleophilic polysubstituted aryl acridinium esters and to novel conjugates thereof. The novel nucleophilic polysubstituted aryl acridinium esters and novel conjugates thereof are useful in biological assays, including novel assays for the determination of Vitamin B.sub.12, folate, cortisol, estradiol, and thromboxane B.sub.2.



09/690,353

L6 ANSWER 47 OF 68 USPATFULL  
 ACCESSION NUMBER: 93:61007 USPATFULL  
 TITLE: Method for the prescreening of drugs targeted to the  
 V3  
 hypervariable loop of the HIV-1 envelope glycoprotein  
 sp 120  
 INVENTOR(S): Neurath, Alexander R., 230 E. 79th St., New York, NY,  
 United States 10021  
 Strick, Nathan, 3243 Lawrence Ave., Oceanside, NY,  
 United States 11572  
 Haberfield, Paul, 1666 52nd St., Brooklyn, NY, United  
 States 11204  
 Jiang, Shibo, 316 W. 95th St. Apr. 525, New York, NY,  
 United States 10025

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 5230998		19930727
APPLICATION INFO.:	US 1991-735640		19910725 (7)
DOCUMENT TYPE:	Utility		
FILE SEGMENT:	Granted		
PRIMARY EXAMINER:	Nucker, Christine M.		
ASSISTANT EXAMINER:	Stucker, Jeffrey		
LEGAL REPRESENTATIVE:	Prishauf, Holtz, Goodman & Woodward		
NUMBER OF CLAIMS:	7		
EXEMPLARY CLAIM:	7		
NUMBER OF DRAWINGS:	19 Drawing Figure(s); 13 Drawing Page(s)		
LINE COUNT:	1592		

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB A method for the rapid screening of a drug targeted to the V3  
 hypervariable loop of the human immunodeficiency virus type 1 or type 2  
 envelope glycoprotein gp 120 comprising measuring the inhibitory effect  
 of the drug on the interaction between gp 120 (or an antigen comprising  
 the V3 hypervariable loop of HIV 1 gp 120 or HIV 2 gp 120) and  
 antibodies specific for the V3 hypervariable loop, and anti-HIV  
 chemotherapy with drugs binding to the V3 hypervariable loop.

L6 ANSWER 48 OF 68 USPATFULL  
 ACCESSION NUMBER: 93:79787 USPATFULL  
 TITLE: Method and composition for double receptor, specific  
 binding assays  
 INVENTOR(S): Bunting, James R., Washington, DC, United States  
 PATENT ASSIGNEE(S): Baxter Diagnostics Inc., Deerfield, IL, United States  
 (U.S. corporation)

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 34394		19930928
APPLICATION INFO.:	US 4271140		19810602 (Original)
	US 1990-629020		19901218 (7)
	US 1978-930130		19780801 (Original)
RELATED APPLN. INFO.:	Continuation-in-part of Ser. No. US 1989-353814, filed on 2 May 1989 And a continuation-in-part of Ser. No.		
US	1978-871478, filed on 23 Jan 1978, now abandoned		
DOCUMENT TYPE:	Reissue		
FILE SEGMENT:	Granted		
PRIMARY EXAMINER:	Nucker, Christine M.		
ASSISTANT EXAMINER:	Preston, David R.		
LEGAL REPRESENTATIVE:	Patterson & Keough		
NUMBER OF CLAIMS:	46		
EXEMPLARY CLAIM:	1,46		
LINE COUNT:	1161		

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The performance of double receptor, specific binding assays is improved  
 by use of a receptor complex having the structure

A.sub.BL (BL).sub.n A.sub.1

wherein BL is a binding ligand, A.sub.BL is a receptor  
 specific for binding ligand, A.sub.1 is a receptor, BL is  
 covalently bonded to A.sub.1 and A.sub.BL is reversibly bonded to BL.  
 Generally A.sub.BL is absorbed onto an insoluble surface and A.sub.1 is  
 an antibody to the substance being assayed. The complex has particular  
 utility in coated tube and rechargeable radioimmunoassay systems.

L6 ANSWER 49 OF 68 CAPLUS COPYRIGHT 2001 ACS  
 ACCESSION NUMBER: 1992:620158 CAPLUS  
 DOCUMENT NUMBER: 117:220158  
 TITLE: A system for enhanced in vivo clearance of diagnostic  
 and/or therapeutic agents by extracorporeal depletion  
 INVENTOR(S): Nilsson, Rune; Lindgren, Lars; Norrgren, Kristina;  
 Sandberg, Bengt; Sjoegren, Hans Olof; Strand, Sven  
 Erik  
 PATENT ASSIGNEE(S): Swed.  
 SOURCE: PCT Int. Appl., 31 pp.  
 CODEN: PIXXD2  
 DOCUMENT TYPE: Patent  
 LANGUAGE: English  
 FAMILY ACC. NUM. COUNT: 1  
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 9212730	A1	19920806	WO 1992-SE20	19920115
W: CA, FI, JP, NO, US				
RW: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LU, MC, NL, SE				
SE 9100142	A	19920718	SE 1991-142	19910117
CA 2100256	AA	19920718	CA 1992-2100256	19920115
EP 567514	A1	19931103	EP 1992-903020	19920115
EP 567514	B1	19930331		
R: AT, BE, CH, DE, DK, ES, FR, GB, IT, LI, LU, NL, SE				
JP 06504542	T2	19940526	JP 1992-503119	19920115
AT 178214	E	19990415	AT 1992-903020	19920115
ES 2130167	T3	19990701	ES 1992-903020	19920115
US 6251394	B1	20010626	US 1993-90047	19931012
PRIORITY APPLN. INFO.:			SE 1991-142	A 19910117
			WO 1992-SE20	W 19920115

AB Non-tissue-bound targeting mols., selective for certain tissues or cells,  
 intended for diagnostic and/or therapeutic applications are removed from  
 blood or reduced to a lower concn. by an extracorporeal system contg.  
 immobilized agents (e.g. receptors) which selectively bind the targeting  
 mols. Targeting mols. are antibodies or their fragments conjugated with  
 cytotoxic drugs, radioisotopes or activators of prodrugs, i.e.  
 enzymes.

L6 ANSWER 50 OF 68 BIOSIS COPYRIGHT 2001 BIOSIS  
 ACCESSION NUMBER: 1993:34784 BIOSIS  
 DOCUMENT NUMBER: PREV199395022984  
 TITLE: IGF-I binding and IGF-I mRNA expression in the  
 post-ischemic regenerating rat kidney.  
 AUTHOR(S): Matejka, Goran L. (1); Jennische, Eva  
 CORPORATE SOURCE: (1) Dep. of Histol., Univ. of Goteborg, Goteborg Sweden  
 SOURCE: Kidney International, (1992) Vol. 42, No. 5, pp.  
 1113-1123.  
 DOCUMENT TYPE: ISSN: 0085-2538.  
 LANGUAGE: Article  
 English

AB The localization of IGF-I peptide and IGF-I mRNA was  
 investigated in the post-ischemic regeneration rat kidney using  
 immunohistochemistry and non-radioactive in situ hybridization  
 techniques. In addition, the distribution and relative quantity of IGF-I  
 binding sites were studied by autoradiographic ligand-binding  
 techniques. Two and three days after the injury, morphological signs of  
 an intense regenerative activity was evident. By this time a substantial  
 number of the regenerating cells were stained with a monoclonal antibody  
 against the M1 subunit of ribonucleotide reductase, a proliferative  
 marker used. Low proliferative tubular cells, replacing those that had been  
 injured, were seen lining the tubular basement membrane. By seven days,  
 the morphology in the cortex was quite normalized while cells of S3  
 segments in the outer medulla remained dedifferentiated. The regenerative  
 cells expressed IGF-I peptide and IGF-I mRNA in a transient  
 manner and this was found to correlate better to cell differentiation  
 than cell division. In addition, non-tubular cells, predominantly macrophages,  
 expressed both the IGF-I peptide and the mRNA. The IGF-I binding  
 was significantly increased in the regenerative zone at all times studied  
 and began to decline at day seven. The binding characteristics were found  
 to be compatible with binding to the IGF-I receptor. Altogether, these  
 findings provide circumstantial evidence that IGF-I is of tropic  
 importance in the regeneration of renal tubular cells. The data are  
 compatible with a local production and action of IGF-I, suggesting an  
 autocrine and/or paracrine mode of action during the regenerative  
 process.

09/690,353

L6 ANSWER 51 OF 68 USPATFULL  
 ACCESSION NUMBER: 91:60741 USPATFULL  
 TITLE: Redox polymerization diagnostic test composition and method for immunoassay and nucleic acid assay  
 INVENTOR(S): Oster, Gerald, 241 W. 11th St., New York, NY, United States 10014  
 Davis, Baruch J., Mount Sinai Medical Center, 1 Gustav Levy Pl., New York, NY, United States 10029

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 5035997		19910730
APPLICATION INFO.:	US 1989-312525		19890217 (7)
DOCUMENT TYPE:	Utility		
FILE SEGMENT:	Granted		
PRIMARY EXAMINER:	Wax, Robert A.		
ASSISTANT EXAMINER:	Marschel, Ardin H.		
LEGAL REPRESENTATIVE:	Sprung Horn Kramer & Woods		
NUMBER OF CLAIMS:	22		
EXEMPLARY CLAIM:	1		
LINE COUNT:	747		

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB A diagnostic test composition for detecting and measuring an analyte possessing biologic activity, the composition comprising

(a) A redox catalyst system capable of converting a monomer to a polymer, the monomer capable of undergoing addition polymerization, the redox catalyst system comprising one or more chemical moieties with

- 1) the analyte comprising at least one such moiety or
- 2) in the case that the analyte lacks a redox catalyst property, the analyte is linked by a specific ligand to at least one such moiety or is linked by the specific ligand to a generator of at least one such moiety, and

(b) at least one monomer capable of undergoing addition polymerization.

L6 ANSWER 52 OF 68 USPATFULL  
 ACCESSION NUMBER: 91:42637 USPATFULL  
 TITLE: Photopolymerization diagnostic test composition and method for immunoassay and nucleic acid assay  
 INVENTOR(S): Oster, Gerald, 241 W. 11th St., New York, NY, United States 10014  
 Davis, Baruch J., Mount Sinai Medical Center, 1 Gustav Levy Pl., New York, NY, United States 10029

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 5019496		19910528
APPLICATION INFO.:	US 1989-312544		19890217 (7)
DOCUMENT TYPE:	Utility		
FILE SEGMENT:	Granted		
PRIMARY EXAMINER:	Wax, Robert A.		
ASSISTANT EXAMINER:	Marschel, Ardin H.		
LEGAL REPRESENTATIVE:	Sprung Horn Kramer & Woods		
NUMBER OF CLAIMS:	40		
EXEMPLARY CLAIM:	1		
LINE COUNT:	1091		

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB A diagnostic test composition for detecting and measuring an analyte possessing biologic activity comprising

(a) a photocatalyst system capable of converting a monomer to a polymer upon exposure to light, the monomer capable of undergoing addition polymerization, the photocatalyst system comprising one or more chemical moieties, with

- (1) the analyte comprising at least one such moiety or generating at least one such moiety or
- (2) in the case that the analyte lacks a photocatalyst property, the analyte is linked by a specific ligand to at least one such moiety or is linked by the specific ligand to a generator of at least one such moiety and

(b) at least one monomer capable of undergoing addition polymerization.

L6 ANSWER 53 OF 68 EMBASE COPYRIGHT 2001 ELSEVIER SCI. B.V.  
 ACCESSION NUMBER: 91108459 EMBASE  
 DOCUMENT NUMBER: 1991108459  
 TITLE: Ligand-exchange radiochromatographic resolution of [3H]DL-valine, [3H]DL-leucine and [3H]DL-methionine using a reverse-phase column in the presence of cupric acetate and GMP or cyanocobalamin.  
 AUTHOR: Fukuhara T.; Isoyama M.; Tanaka M.; Yuasa S.  
 CORPORATE SOURCE: Department of Biology, College of General Education, Osaka University, Osaka 560, Japan  
 SOURCE: Applied Radiation and Isotopes, (1991) 42/5 (457-462). ISSN: 0883-2889 CODEN: ARISEF  
 COUNTRY: United Kingdom  
 DOCUMENT TYPE: Journal; Article  
 FILE SEGMENT: 023 Nuclear Medicine  
 029 Clinical Biochemistry  
 LANGUAGE: English  
 SUMMARY LANGUAGE: English

AB [3H]DL-valine, [3H]DL-leucine and [3H]DL-methionine were resolved using ligand-exchange chromatography (reverse-phase) in the presence of cupric acetate [Cu(II)], and 5'-guanosine monophosphate or cyanocobalamin.  
 The assignment of the resolved enantiomers was carried out by means of co-chromatography with non-labeled DL-amino acids after modifying them with fluorodinitrobenzene. The optical purity of the enantiomers was estimated to be greater than 99%. The resolved enantiomers were subjected to bioassay, which showed that the enantiomers were biochemically active.

L6 ANSWER 54 OF 68 USPATFULL  
 ACCESSION NUMBER: 90:54554 USPATFULL  
 TITLE: Assay for sulfhydryl amino acids and methods for detecting and distinguishing cobalamin and folic acid deficiency  
 INVENTOR(S): Allen, Robert H., Englewood, CO, United States  
 Stabler, Sally P., Denver, CO, United States  
 Lindenbaum, John, New York, NY, United States  
 PATENT ASSIGNEE(S): University Patents, Inc., Westport, CT, United States (U.S. corporation)

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 4940658		19900710
APPLICATION INFO.:	US 1986-933553		19861120 (6)
DOCUMENT TYPE:	Utility		
FILE SEGMENT:	Granted		
PRIMARY EXAMINER:	Kepplinger, Esther M.		
ASSISTANT EXAMINER:	Scheiner, Toni R.		
LEGAL REPRESENTATIVE:	Yahwak & Associates		
NUMBER OF CLAIMS:	34		
EXEMPLARY CLAIM:	18		
NUMBER OF DRAWINGS:	7 Drawing Figure(s); 7 Drawing Page(s)		
LINE COUNT:	2375		

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB Method for determining levels of sulfhydryl amino acids, particularly total homocysteine levels in samples of body tissue from warm-blooded animals, methods of detecting cobalamin and folic acid deficiency using an assay for total homocysteine levels, and methods for distinguishing cobalamin from folic acid deficiency using an assay for total homocysteine levels in conjunction with an assay for methylmalonic acid.

09/690,353

L6 ANSWER 55 OF 68 USPATFULL  
 ACCESSION NUMBER: 90:15471 USPATFULL  
 TITLE: Cascade immunoassay by multiple binding reactions  
 INVENTOR(S): Mapes, James P., Raleigh, NC, United States  
 Hoke, Randal A., Cary, NC, United States  
 PATENT ASSIGNEE(S): Becton, Dickinson and Company, Franklin Lakes, NJ, United States (U.S. corporation)

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 4904583		19900227
APPLICATION INFO.:	US 1987-53896		19870526 (7)
DOCUMENT TYPE:	Utility		
FILE SEGMENT:	Granted		
PRIMARY EXAMINER:	Warden, Robert J.		
ASSISTANT EXAMINER:	Spiegel, Jack		
LEGAL REPRESENTATIVE:	Brown, Richard E.		
NUMBER OF CLAIMS:	31		
EXEMPLARY CLAIM:	1		
NUMBER OF DRAWINGS:	8 Drawing Figure(s); 7 Drawing Page(s)		
LINE COUNT:	785		

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB A method for enzyme immunoassay includes contacting under binding conditions a liquid suspected of containing an analyte, an antianalyte affixed to a solid support and a tracer having an enzyme conjugated thereto. A bound fraction is separated from the liquid and incubated in a second liquid with a masked ligand. The masked ligand is converted by the enzyme on the bound fraction to give free ligand which binds to an antiligand. A signal system, such as a signal enzyme and substrate therefor, or a label-loaded vesicle and vesicle lysing agent, is added to generate a signal used to detect or measure the analyte in the liquid. The invention includes a kit of materials useful in performing the assay of the invention.

L6 ANSWER 56 OF 68 USPATFULL  
 ACCESSION NUMBER: 87:69949 USPATFULL  
 TITLE: Enzymatic reactions using magnetic particles  
 INVENTOR(S): Whitehead, Roy A., Hingham, MA, United States  
 Chagnon, Mark S., Lowell, MA, United States  
 Groman, Ernest V., Brookline, MA, United States  
 PATENT ASSIGNEE(S): Josephson, Lee, Arlington, MA, United States  
 Advanced Magnetica, Inc., Cambridge, MA, United States (U.S. corporation)

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 4698302		19871006
APPLICATION INFO.:	US 1985-744457		19850613 (6)
RELATED APPLN. INFO.:	Division of Ser. No. US 1983-493991, filed on 12 May 1983, now patented, Pat. No. US 4554088		
DOCUMENT TYPE:	Utility		
FILE SEGMENT:	Granted		
PRIMARY EXAMINER:	Foelak, Morton		
ASSISTANT EXAMINER:	Nutter, Nathan M.		
LEGAL REPRESENTATIVE:	Pennie & Edmonds		
NUMBER OF CLAIMS:	16		
EXEMPLARY CLAIM:	1		
NUMBER OF DRAWINGS:	2 Drawing Figure(s); 2 Drawing Page(s)		
LINE COUNT:	1464		

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB A process is provided for the preparation of magnetic particles to which

a wide variety of molecules may be coupled. The magnetic particles can be dispersed in aqueous media without rapid settling and conveniently reclaimed from media with a magnetic field. Preferred particles do not become magnetic after application of a magnetic field and can be redispersed and reused. The magnetic particles are useful in biological systems involving separations.

L6 ANSWER 57 OF 68 USPATFULL  
 ACCESSION NUMBER: 87:66802 USPATFULL  
 TITLE: Magnetic particles for use in separations  
 INVENTOR(S): Chagnon, Mark S., Lowell, MA, United States  
 Groman, Ernest V., Brookline, MA, United States  
 Whitehead, Roy A., Hingham, MA, United States  
 PATENT ASSIGNEE(S): Josephson, Lee, Arlington, MA, United States  
 Advanced Magnetica Inc., Cambridge, MA, United States (U.S. corporation)

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 4695392		19870922
APPLICATION INFO.:	US 1985-744435		19850613 (6)
RELATED APPLN. INFO.:	Division of Ser. No. US 1983-493991, filed on 12 May 1983, now patented, Pat. No. US 4554088		
DOCUMENT TYPE:	Utility		
FILE SEGMENT:	Granted		
PRIMARY EXAMINER:	Foelak, Morton		
ASSISTANT EXAMINER:	Nutter, Nathan M.		
LEGAL REPRESENTATIVE:	Pennie & Edmonds		
NUMBER OF CLAIMS:	12		
EXEMPLARY CLAIM:	1		
NUMBER OF DRAWINGS:	2 Drawing Figure(s); 2 Drawing Page(s)		
LINE COUNT:	1514		

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB A process is provided for the preparation of magnetic particles to which

a wide variety of molecules may be coupled. The magnetic particles can be dispersed in aqueous media without rapid settling and conveniently reclaimed from media with a magnetic field. Preferred particles do not become magnetic after application of a magnetic field and can be redispersed and reused. The magnetic particles are useful in biological systems involving separations.

L6 ANSWER 58 OF 68 USPATFULL  
 ACCESSION NUMBER: 87:66801 USPATFULL  
 TITLE: Magnetic particles for use in separations  
 INVENTOR(S): Whitehead, Roy A., Hingham, MA, United States  
 Chagnon, Mark S., Lowell, MA, United States  
 Groman, Ernest V., Brookline, MA, United States  
 PATENT ASSIGNEE(S): Josephson, Lee, Arlington, MA, United States  
 Advanced Magnetica Inc., Cambridge, MA, United States (U.S. corporation)

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 4695392		19870922
APPLICATION INFO.:	US 1985-744434		19850613 (6)
RELATED APPLN. INFO.:	Division of Ser. No. US 1983-493991, filed on 12 May 1983, now patented, Pat. No. US 4554088		
DOCUMENT TYPE:	Utility		
FILE SEGMENT:	Granted		
PRIMARY EXAMINER:	Kight, John		
ASSISTANT EXAMINER:	Nutter, Nathan M.		
LEGAL REPRESENTATIVE:	Pennie & Edmonds		
NUMBER OF CLAIMS:	11		
EXEMPLARY CLAIM:	1		
NUMBER OF DRAWINGS:	2 Drawing Figure(s); 2 Drawing Page(s)		
LINE COUNT:	1459		

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB A process is provided for the preparation of magnetic particles to which

a wide variety of molecules may be coupled. The magnetic particles can be dispersed in aqueous media without rapid settling and conveniently reclaimed from media with a magnetic field. Preferred particles do not become magnetic after application of a magnetic field and can be redispersed and reused. The magnetic particles are useful in biological systems involving separations.

09/690,353

L6 ANSWER 59 OF 68 USPATFULL  
 ACCESSION NUMBER: 87:41600 USPATFULL  
 TITLE: Magnetic particles for use in separations  
 INVENTOR(S): Josephson, Lee, Arlington, MA, United States  
 PATENT ASSIGNEE(S): Advanced Magnetice, Inc., Cambridge, MA, United States (U.S. corporation)

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 4672040		19870609
APPLICATION INFO.:	US 1985-749692		19850628 (6)
DISCLAIMER DATE:	20021119		

RELATED APPLN. INFO.: Continuation-in-part of Ser. No. US 1983-493991, filed on 12 May 1983, now patented, Pat. No. US 4554088 And Ser. No. US 1985-744351, filed on 13 Jun 1985, now patented, Pat. No. US 4628037 And Ser. No. US 1985-744435, filed on 13 Jun 1985 And Ser. No. US 1985-744434, filed on 13 Jun 1985 And Ser. No. US 1985-744457, filed on 13 Jun 1985

DOCUMENT TYPE: Utility  
 FILE SEGMENT: Granted  
 PRIMARY EXAMINER: Nucker, Christine M.  
 ASSISTANT EXAMINER: Wiedner, Stephen C.  
 LEGAL REPRESENTATIVE: Pennie & Edmonds  
 NUMBER OF CLAIMS: 23  
 EXEMPLARY CLAIM: 1  
 NUMBER OF DRAWINGS: 2 Drawing Figure(s); 2 Drawing Page(s)  
 LINE COUNT: 1770

CAS INDEXING IS AVAILABLE FOR THIS PATENT.  
 AB Methods are provided for the use of magnetically responsive particles in

and systems in which the separation of certain molecules, macromolecules and cells from the surrounding medium is desirable. The magnetically responsive particles may be coupled to a wide variety of molecules. The magnetic particles can be dispersed in aqueous media without rapid settling and conveniently reclaimed from media with a magnetic field. Preferred particles do not become magnetic after application of a magnetic field and can be redispersed and reused.

L6 ANSWER 61 OF 68 USPATFULL  
 ACCESSION NUMBER: 86:69734 USPATFULL  
 TITLE: Binding assays employing magnetic particles  
 INVENTOR(S): Chagnon, Mark S., Lowell, MA, United States  
 Groman, Ernest V., Brookline, MA, United States  
 Josephson, Lee, Arlington, MA, United States  
 Whitehead, Roy A., Hingham, MA, United States  
 PATENT ASSIGNEE(S): Advanced Magnetice, Inc., Cambridge, MA, United States (U.S. corporation)

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 4628037		19861209
APPLICATION INFO.:	US 1985-744351		19850613 (6)
RELATED APPLN. INFO.:	Division of Ser. No. US 1983-493991, filed on 12 May 1983, now patented, Pat. No. US 4554488		

DOCUMENT TYPE: Utility  
 FILE SEGMENT: Granted  
 PRIMARY EXAMINER: Nucker, Christine M.  
 ASSISTANT EXAMINER: Wiedner, Stephen C.  
 LEGAL REPRESENTATIVE: Pennie & Edmonds  
 NUMBER OF CLAIMS: 11  
 EXEMPLARY CLAIM: 1  
 NUMBER OF DRAWINGS: 2 Drawing Figure(s); 2 Drawing Page(s)  
 LINE COUNT: 1495

CAS INDEXING IS AVAILABLE FOR THIS PATENT.  
 AB A process is provided for the preparation of magnetic particles to which a wide variety of molecules may be coupled. The magnetic particles can be dispersed in aqueous media without rapid settling and conveniently reclaimed from media with a magnetic field. Preferred particles do not become magnetic after application of a magnetic field and can be redispersed and reused. The magnetic particles are useful in biological systems involving separations.

L6 ANSWER 60 OF 68 USPATFULL  
 ACCESSION NUMBER: 87:41588 USPATFULL  
 TITLE: Compositions and method for simultaneous multiple array  
 of analytes using radioisotope chelate labels  
 INVENTOR(S): Olson, Douglas R., Doylestown, PA, United States  
 PATENT ASSIGNEE(S): ICN Micromedic Systems, Inc., Costa Mesa, CA, United States (U.S. corporation)

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 4672028		19870609
APPLICATION INFO.:	US 1984-612979		19840523 (6)

DOCUMENT TYPE: Utility  
 FILE SEGMENT: Granted  
 PRIMARY EXAMINER: Nucker, Christine M.  
 LEGAL REPRESENTATIVE: Lyon & Lyon  
 NUMBER OF CLAIMS: 47  
 EXEMPLARY CLAIM: 1  
 LINE COUNT: 784

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB Compounds useful in a simultaneous multiple assay for analytes such as steroids, proteins, peptides, carbohydrates or drugs. The compound or compounds are prepared by labelling an individual analyte with a radioisotope through a chelating agent to form a coordinated compound. The assay uses one or more chelated labelled analytes with one or more labelled analytes wherein each radioisotope is different.

L6 ANSWER 62 OF 68 CAPLUS COPYRIGHT 2001 ACS  
 ACCESSION NUMBER: 1986:84931 CAPLUS  
 DOCUMENT NUMBER: 104:84931  
 TITLE: Simultaneous multiple assays and compounds and compositions useful in them  
 INVENTOR(S): Olson, Douglas Richard  
 PATENT ASSIGNEE(S): Micromedic Systems, Inc., USA  
 SOURCE: Eur. Pat. Appl., 26 pp.  
 CODEN: EFXKDW  
 DOCUMENT TYPE: Patent  
 LANGUAGE: English  
 FAMILY ACC. NUM. COUNT: 2  
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
EP 165716	A1	19851227	EP 1985-303564	19850521
EP 165716	B1	19900131		
R: AT, BE, CH, DE, FR, GB, IT, LI, LU, NL, SE				
US 4672028	A	19870609	US 1984-612979	19840523
AT 50066	E	19900215	AT 1985-303564	19850521
AU 8542798	A1	19851128	AU 1985-42798	19850523
AU 582970	B2	19890413		
JP 6100092	A2	19860106	JP 1985-111312	19850523
PRIORITY APPLN. INFO.:			US 1984-612979	19840523
			EP 1985-303564	19850521

AB Simultaneous multiple assays for org. species (e.g., steroids, proteins, peptides, carbohydrates, drugs) are described. This procedure makes use of coordinated compds., which are prepd. by labeling an individual org. species with a radioisotope through a chelating agent of general structure (HO<sub>2</sub>CCH<sub>2</sub>)<sub>2</sub>NCHR[CH<sub>2</sub>N(CH<sub>2</sub>CO<sub>2</sub>H)C H<sub>2</sub>]nCH<sub>2</sub>N(CH<sub>2</sub>CO<sub>2</sub>H)<sub>2</sub> where R = Ph, or Ph substituted with NO<sub>2</sub>, NH<sub>2</sub>, and/or SO<sub>3</sub>H and n = 0 or 1. Radiolabeled org. species compd. individually distinguishable radionuclides are combined in a variety of configurations to measure more org. species simultaneously using radioassay. For example, for simultaneous detn. of LH and FSH, 57Co-labeled LH was prepd. by mixing lyophilized LH with diethylenetriaminepentaacetic anhydride (I) in Na<sub>2</sub>CO<sub>3</sub> soln. The resulting LH-I was mixed with 57CoCl<sub>2</sub> in HCl to yield 57Co-labeled LH. 125I-labeled FSH was prepd. by mixing FSH antigen with Na<sub>2</sub>125I in phosphate-buffered saline. A test sample or std. was mixed with antiserum and incubated for 1 h at 37.degree.. Tracer solns. were added and the reaction mixt. was further incubated for 1 h at room temp. After addn. of pptg. soln., the mixt. was incubated for 10 min at room temp. and centrifuged. The liq. was discarded and the tubes were counted for 125I and 57Co in a gamma-counter. Unknown samples were read from a std. curve.

09/690,353

L6 ANSWER 63 OF 68 USPATFULL  
 ACCESSION NUMBER: 85:68137 USPATFULL  
 TITLE: Magnetic particles for use in separations  
 INVENTOR(S): Whitehead, Roy A., Hingham, MA, United States  
 Chagnon, Mark S., Lowell, MA, United States  
 Groman, Ernest V., Brookline, MA, United States  
 Josephson, Lee, Arlington, MA, United States  
 PATENT ASSIGNEE(S): Advanced Magnetics Inc., Cambridge, MA, United States  
 (U.S. corporation)

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 4554088		19851119
APPLICATION INFO.:	US 1983-493991		19830512 (6)
DOCUMENT TYPE:	Utility		
FILE SEGMENT:	Granted		
PRIMARY EXAMINER:	Demers, Arthur P.		
LEGAL REPRESENTATIVE:	Pennie & Edmonds		
NUMBER OF CLAIMS:	11		
EXEMPLARY CLAIM:	1		
NUMBER OF DRAWINGS:	2 Drawing Figure(s); 2 Drawing Page(s)		
LINE COUNT:	1501		

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB A process is provided for the preparation of magnetic particles to which

a wide variety of molecules may be coupled. The magnetic particles can be dispersed in aqueous media without rapid settling and conveniently reclaimed from media with a magnetic field. Preferred particles do not become magnetic after application of a magnetic field and can be redispersed and reused. The magnetic particles are useful in biological systems involving separations.

L6 ANSWER 64 OF 68 USPATFULL  
 ACCESSION NUMBER: 84:25942 USPATFULL  
 TITLE: Detecting intrinsic factor blocking site antibody  
 INVENTOR(S): Ellis, James E., Stoughton, MA, United States  
 Lidgard, Graham P., Wellesley, MA, United States  
 Odstrchel, Gerald, Walpole, MA, United States  
 Riceberg, Louis J., Needham, MA, United States  
 PATENT ASSIGNEE(S): Corning Glass Works, Corning, NY, United States (U.S. corporation)

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 4447528		19840508
APPLICATION INFO.:	US 1981-291354		19810810 (6)
DOCUMENT TYPE:	Utility		
FILE SEGMENT:	Granted		
PRIMARY EXAMINER:	Marantz, Sidney		
LEGAL REPRESENTATIVE:	Maycock, W. E.		
NUMBER OF CLAIMS:	7		
EXEMPLARY CLAIM:	1		
LINE COUNT:	395		

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB A radioassay procedure and reagent kit therefor for detecting auto blocking antibody, such as auto blocking antibody which interferes with the complexation of intrinsic factor with vitamin B.sub.12. A receptor, i.e., intrinsic factor, is immobilized on a support and the amount of ligand, i.e., vitamin B.sub.12, capable of binding therewith in the presence of a biological fluid sample is determined.

L6 ANSWER 65 OF 68 USPATFULL  
 ACCESSION NUMBER: 81:30332 USPATFULL  
 TITLE: Method and composition for double receptor, specific binding assays  
 INVENTOR(S): Bunting, James R., Washington, DC, United States  
 PATENT ASSIGNEE(S): Baxter Travenol Laboratories, Inc., Deerfield, IL, United States (U.S. corporation)

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 4271140		19810602
APPLICATION INFO.:	US 1978-930130		19780801 (5)
RELATED APPLN. INFO.:	Continuation-in-part of Ser. No. US 1978-871478, filed on 23 Jan 1978, now abandoned		
DOCUMENT TYPE:	Utility		
FILE SEGMENT:	Granted		
PRIMARY EXAMINER:	Padgett, Benjamin R.		
ASSISTANT EXAMINER:	Nucker, Christine M.		
LEGAL REPRESENTATIVE:	Plattery, Paul C., Flynn, Lawrence W., Hensley, Max D.		
NUMBER OF CLAIMS:	45		
EXEMPLARY CLAIM:	1		
LINE COUNT:	1050		

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The performance of double receptor, specific binding assays is improved by use of a receptor complex having the structure

A.sub.BL (BL).sub.n A.sub.1

wherein BL is a binding ligand, A.sub.BL is a receptor specific for binding ligand, A.sub.1 is a receptor, BL is covalently bonded to A.sub.1 and A.sub.BL is reversibly bonded to BL. Generally A.sub.BL is adsorbed onto an insoluble surface and A.sub.1 is an antibody to the substance being assayed. The complex has particular utility in coated tube and rechargeable radioimmunoassay systems.

L6 ANSWER 66 OF 68 USPATFULL  
 ACCESSION NUMBER: 81:27515 USPATFULL  
 TITLE: Automated direct serum radioassay  
 INVENTOR(S): Reese, Max, Salt Lake City, UT, United States  
 PATENT ASSIGNEE(S): Becton Dickinson & Company, Paramus, NJ, United States (U.S. corporation)

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 4268494		19810519
APPLICATION INFO.:	US 1978-920801		19780630 (5)
DISCLAIMER DATE:	19950822		
RELATED APPLN. INFO.:	Continuation-in-part of Ser. No. US 1977-774390, filed on 4 Mar 1977, now patented, Pat. No. US 4108976		
DOCUMENT TYPE:	Utility		
FILE SEGMENT:	Granted		
PRIMARY EXAMINER:	Schafer, Richard E.		
ASSISTANT EXAMINER:	Nucker, Christine M.		
LEGAL REPRESENTATIVE:	Marn, Louis E., Olstein, Elliot M.		
NUMBER OF CLAIMS:	7		
EXEMPLARY CLAIM:	1		
LINE COUNT:	499		

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB Automated radioassay in which a serum is diluted and pre-incubated in the presence of a radioactive labeled ligand, such as an antigen, and a binder, such as an antibody, specific to the ligand. The pre-incubated mixture is flowed through a chamber containing binder specific to the ligand supported on a solid support, and the labeled and unlabeled ligand not bound to the binder in pre-incubation are bound to the receptor on flow through the chamber. An eluting solution is flowed through the chamber to release the ligand bound to the binder in the chamber for reuse thereof. By counting the radioactivity of one or both of the fraction which flows through the chamber or which is subsequently released therefrom the quantity of a specific ligand in the serum may be assayed.

09/690,353

L6 ANSWER 67 OF 68 USPATFULL  
 ACCESSION NUMBER: 78:52922 USPATFULL  
 TITLE: in vitro diagnostic test  
 INVENTOR(S): Ithakisios, Dionysis S., St. Paul, MN, United States  
 PATENT ASSIGNEE(S): Minnesota Mining and Manufacturing Company, St. Paul, MN, United States (U.S. corporation)

NUMBER	KIND	DATE
US 4115534		19780919
US 1977-808339		19770620 (5)

Continuation-in-part of Ser. No. US 1976-715933, filed on 19 Aug 1976, now abandoned

DOCUMENT TYPE: Utility  
 FILE SEGMENT: Granted  
 PRIMARY EXAMINER: Padgett, Benjamin R.  
 ASSISTANT EXAMINER: Nucker, Christine M.  
 LEGAL REPRESENTATIVE: Alexander, C., Sell, Donald M., Lilly, James V.  
 NUMBER OF CLAIMS: 25  
 EXEMPLARY CLAIM: 1  
 NUMBER OF DRAWINGS: 2 Drawing Figure(s); 1 Drawing Page(s)  
 LINE COUNT: 804

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB A method for determining the concentration of substances in biological fluids (e.g., drugs, hormones, vitamins and enzymes) is disclosed wherein magnetically responsive, permeable, solid, water-insoluble microparticles are employed.

L6 ANSWER 68 OF 68 CAPLUS COPYRIGHT 2001 ACS  
 ACCESSION NUMBER: 1978:34141 CAPLUS  
 DOCUMENT NUMBER: 88:34141  
 TITLE: Detection of genetic variation with radioactive ligands. I.  
 Electrophoretic screening of plasma proteins with a selected panel of compounds  
 Cavalli-Sforza, Luigi L.; Daiger, Stephen P.; Rummel, Diane P.  
 AUTHOR(S): Dep. Genet., Stanford Univ. Med. Cent., Stanford, Calif., USA  
 CORPORATE SOURCE: Am. J. Hum. Genet. (1977), 29(6), 581-92  
 SOURCE: CODEN: AJHGAG  
 DOCUMENT TYPE: Journal  
 LANGUAGE: English

AB To detect new genetic variation in human plasma proteins, a panel of 63 radioactive substances were screened as potential radioligands using polyacrylamide gel electrophoresis and autoradiog. Vitamins, hormones, drugs, amino acids, purines, pyrimidines, sugars, and lipids labeled with <sup>14</sup>C or other radionuclides were among those substances tested. A majority bound to albumin and a smaller fraction to prealbumins and lipoproteins. Several vitamins and hormones bound to specific .alpha. and .beta. globulins. Electrophoretic polymorphisms of the vitamin D-binding protein (group-specific component), a vitamin B12-binding protein (transcobalamin II), and thyroxine-binding .alpha. globulin were obsd. Testosterone-binding .beta. globulin showed an electrophoretic polymorphism in Caucasians and a possible deficiency allele. Transcortin showed an electrophoretic doublet in all persons tested but no electrophoretic variation. A protein binding a deriv. of norepinephrine or epinephrine was identified as transferrin. A nonpolymorphic protein running cathodal to albumin and binding a deriv. of riboflavin was tentatively identified as a fraction of albumin with mobility altered as a result of interaction with the ligand.